Access DB#	

SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: Art Unit: Mail Box and Bldg/Room Location	lumber 30 <u>8 - 469 (</u>		09/781645
If more than one search is subm		e searches in order of need.	*****
Please provide a detailed statement of the Include the elected species or structures, k utility of the invention. Define any terms known. Please attach a copy of the cover s	eywords, synonyms, acron that may have a special me heet, pertinent claims, and	yms, and registry numbers, and combinaning. Give examples or relevant citat abstract.	ne with the concept or iions, authors, etc, if
Title of Invention: Induc	ing Produc	tion of Isofla	vone to prent
Inventors (please provide full names):	Jevrence (Fraham; Lian-1	Mei Graham;
	Sevena L	andini	· · .
Earliest Priority Filing Date:			
For Sequence Searches Only Please include appropriate serial number.	le all pertinent information (j	parent, child, divisional, or issued patent n	numbers) along with the
Search claim	701 100	crop or stem of Loliage	r rod ur
Search	- gant or	Loliage	e e e e e e e e e e e e e e e e e e e
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. Slard	Compa	A. (5)	
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Search Claim	3022	Jors op orthou	anadate,
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STAFF USE ONLY Searcher:	Type of Search NA Sequence (#)	Vendors and cost where a	pplicable
Searcher Phone #:	AA Sequence (#)	Dialog	
Searcher Location:	Structure (#)	Questel/Orbit	
Date Searcher Picked Up: 8/22	Bibliographic	Dr.Link	·
Date Completed: 8/22	Litigation	Lexis/Nexis	·
Searcher Prep & Review Time: 70	Fulltext	Sequence Systems	<u> </u>
Clerical Prep Time:	Patent Family	WWW/Internet	. , , ,
Online Time:	Other	Other (specify)	

cités cover claim 1 (isoflavone induction) but these are plant cultures (not a PRYOR 09/781,695 plant) parket # 2

=> d ibib abs hitstr ind 195 1-6

L95 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: DOCUMENT NUMBER:

2002:440428 HCAPLUS

137:244656

TITLE:

Phenolic antioxidant compounds produced by in vitro

shoots of sage (Salvia officinalis L.)

AUTHOR(S):

Santos-Gomes, Paula C.; Seabra, Rosa M.; Andrade,

Paula B.; Fernandes-Ferreira, Manuel

CORPORATE SOURCE:

Department of Biology, University of Minho, Braga, 4710-057, Port.

SOURCE:

Plant Science (Shannon, Ireland) (2002), 162(6),

981-987

CODEN: PLSCE4; ISSN: 0168-9452 Elsevier Science Ireland Ltd.

PUBLISHER: DOCUMENT TYPE:

English LANGUAGE:

In vitro shoots of sage (Salvia officinalis L.) were established under four different cytokinin supplementations by culturing nodal segments excised from aseptically germinated seedlings. The highest rates of shoot proliferation and linear shoot growth occurred with the supplementation of 1.5 mg/l benzyladenine and 0.05 mg/l dichlorophenoxyacetic acid. However, under these conditions, the specific prodn. of total antioxidant phenolics was the lowest. Variation in kinetin (KIN) concn. (1.5; 2.0; 4.0 mg/l), in the presence of 0.05 mg/l 2,4-D, did not influence significantly the rates of shoot proliferation and linear shoot growth but influenced the prodn. of antioxidant phenolics and biomass. Seventeen compds. were identified in the antioxidant phenolic exts. from shoots: gallic acid, 3-O-caffeoylquinic acid, 5-O-caffeoylquinic acid, caffeic acid, and rosmarinic acid, as phenolic acids; hesperetin, apigenin, hispidulin, cirsimaritin, and genkwanin, as flavonoids; epirosmanol, epirosmanol Me ether, carnosol, epiisorosmanol Et ether, rosmadial, carnosic acid, and Me carnosate, as phenolic diterpenes. With exception of carnosic acid and Me carnosate, all the other phenolic compds. were also identified in a com. sample of this species. Rosmarinic acid and carnosol were the main compds. in all the antioxidant phenolic exts. The increase in concn. of KIN decreased the accumulation of the most of phenolic diterpenes, particularly that of carnosol.

94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations)

RN 94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN

11-1 (Plant Biochemistry) CC Section cross-reference(s): 16, 17

phenolic sage antioxidant; Salvia phenolic antioxidant

ST Antioxidants

Sage (Salvia officinalis)

(phenolic antioxidant compds. produced by in vitro shoots of sage)

IT Plant tissue culture

(phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations)

IT Flavon ids

RL: BSU (Biological study, unclassified); BIOL (Biological study)

(phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations) Cytokinins IT RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations) Carboxylic acids, biological studies RL: BSU (Biological study, unclassified); BIOL (Biological study) (phenolic; phenolic antioxidant compds. produced by in vitro shoots of sage) Growth and development, plant IT (shoot formation; phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations) 149-91-7, Gallic acid, biological studies 327-97-9, 3-0-Caffeoylquinic acid 331-39-5, Caffeic acid 437-64-9, Genkwanin 5520-36-5, Apigenin 906-33-2, 5-0-Caffeoylquinic acid 520-33-2, Hesperetin 1447-88-7 3650-09-7, Carnosic acid 5957-80-2, Carnosol 20283-92-5, Rosmarinic acid _24703-38-6, Epirosmanol Cirsimaritin methyl ether 82684-06-8, Methyl carnosate 85514-31-4, Rosmadial 177027-96-2, Epiisorosmanol ethyl ether 93380-12-2, Epirosmanol RL: BSU (Biological study, unclassified); BIOL (Biological study) (phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations) 94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies 1214-39-7, Benzyladenine 525-79-1, Kinetin RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (phenolic antioxidant compds. produced by in vitro shoots of sage under different cytokinin supplementations) REFERENCE COUNT: 31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L95 ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN 2002:419531 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 137:168317 Obtaining flavonoids by using tissue culture anther TITLE: and style of Flos sophorae Wang, Lijuan; Li, Feng; Yang, Jianxiong School of Sciences, Xi'an Jiaotong University, Xi'an, AUTHOR(S): CORPORATE SOURCE: 710049, Peop. Rep. China Xi'an Jiaotong Daxue Xuebao (2002), 36(3), 328-330 SOURCE: CODEN: HCTPDW; ISSN: 0253-987X PUBLISHER: Xi'an Jiaotong Daxue Xuebao Bianjibu DOCUMENT TYPE: Journal LANGUAGE: Chinese Flavonoids contg. rutin is manufd. by tissue culture of style and anther of the flower of Sophora (huaimi). Induction of callus and prodn. of flavonoids are improved with the addn. of hormones such as 2.4-D and 6-BA. 94-75-7, 2,4-D, biological studies RL: BSU (Biological study, unclassified); BIOL (Biological study) (obtaining flavonoids by using tissue culture anther and style of Flos sophorae) 94-75-7 HCAPLUS RN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) ČN

```
16-2 (Fermentation and Bioindustrial Chemistry)
     rutin flavonoid manuf Sophora tissue culture
ST
IT
     Plant tissue culture
        (callus; obtaining flavonoids by using tissue culture anther and style
        of Flos sophorae)
     Natural products, pharmaceutical
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (huaimi; obtaining flavonoids by using tissue culture anther
        and style of Flos sophorae)
IT
     Sophora
        (obtaining flavonoids by using tissue culture anther and style of Flos
        sophorae)
     Flavonoids
IT
     RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP
     (Preparation)
        (obtaining flavonoids by using tissue culture anther and style of Flos
        sophorae)
IT
     Hormones, plant
     RL: BSU (Biological study, unclassified); BTOL (Biological study)
        (obtaining flavonoids by using tissue culture anther and style of Flos
        sophorae)
     153-18-4P, Rutin
IT
     RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP
     (Preparation)
        (obtaining flavonoids by using tissue culture anther and style of Flos
        sophorae)
     94-75-7, 2,4-D, biological studies 1214-39-7, 6-BA
RL: BSU (Biological study, unclassified); BIOL (Biological study)
TT
        (obtaining flavonoids by using tissue culture anther and style of Flos
        sophorae)
L95 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN
                         2001:625766 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         135:315972
                         Growth and flavonoid production in
TITLE:
                         Bellis perennis L. callus cultures
AUTHOR(S):
                          Siatka, T.; Kasparova, M.
                         Department of Pharmacognosy, Charles University,
CORPORATE SOURCE:
                          Hradec Kralove, 500 05, Czech Rep.
                          Herba Polonica (2001), 47(1), 17-21
SOURCE:
                          CODEN: HPBIA9; ISSN: 0018-0599
                         Instytut Roslin i Przetworow Zielarskich
PUBLISHER:
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English.
     Effects of a cytokinin benzyladenine (at the concns. of 0.1 or 1 mg/l) in
     combination with auxins 2,4-dichlorophenoxyacetic acid,
     .alpha.-naphthaleneacetic acid or .beta.-indoleacetic acid (at the concns.
     of 0.1; 1 or 10 mg/l) on the cell growth and flavonoid
     prodn. in Bellis perennis callus cultures were studied.
                                                               The best
     results were obtained by combination of 0.1 mg/l 2,4-dichlorophenoxyacetic
     acid with 0.1 mg/l benzyladenine for the growth and by combination of 1
     mg/l .alpha.-naphthaleneacetic acid with 0.1 mg/l benzyladenine for the
     flavonoid prodn.
     94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies
IT
     RL: BAC (Biological activity or effector,-except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (growth and flavonoid prodn. in Bellis perennis
        callus cultures)
RN
     94-75-7 HCAPLUS
     Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
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0- CH2- CO2H
     11-3 (Plant Biochemistry)
     Section cross-reference(s): 16
     Bellis callus culture flavonoid prodn phytohormone
ST
IT
     Plant tissue culture
        (callus; growth and flavonoid prodn. in Bellis
        perennis callus cultures)
IT
     Bellis perennis
```

(growth and flavonoid prodn. in Bellis perennis callus cultures)

IT Auxins

Hormones, plant

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (growth and flavonoid prodn. in Bellis perennis callus cultures)

IT Flavonoids

RL: BMF (Bioindustrial manufacture); BIOL (Biological study); PREP (Preparation)

(growth and flavonoid prodn. in Bellis perennis

callus cultures)

86-87-3, .alpha.-Naphthaleneacetic acid 87-51-4, .beta.-Indoleacetic acid, biological studies 94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies 1214-39-7, BA RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (growth and flavonoid prodn. in Bellis perennis

callus cultures)

REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L95 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

2001:621314 HCAPLUS

DOCUMENT NUMBER:

135:329300

TITLE:

Promotive effect of auxins on UDP-glucose:flavonol

glucosyltransferase activity in Vitis sp. cell

AUTHOR(S):

Kokubo, Tetsuro; Ambe-Ono, Yukiko; Nakamura, Mayumi;

CORPORATE SOURCE:

Ishida, Hidekatsu; Yamakawa, Takashi; Kodama, Tohru Department of Biotechnology, The University of Tokyo,

Tokyo, 113-8657, Japan

SOURCE:

Journal of Bioscience and Bioengineering (2001),

91(6), 564-569 CODEN: JBBIF6; ISSN: 1389-1723

PUBLISHER:

Society for Bioscience and Bioengineering, Japan

DOCUMENT TYPE: Journal

English LANGUAGE:

The addn. of 2,4-dichlorophenoxyacetic acid (2, 4-D) to Vitis sp. cell cultures significantly enhanced the prodn. of quercetin 3,7,4'-tri-O-glucoside, 3,7-di-O-glucoside and 3,4'-di-O-glucoside from quercetin. This enhancement of glucosylation by 2,4-D was also obsd. in cell cultures of other plant species. The activity of UDP-glucose:flavonol glucosyltransferase (UFGT) in cell-free exts. of Vitis sp. cell cultures increased approx. 10-fold, 48 h after the addn. of 2,4-D to the culture medium. The UFGT activity increased linearly up to 15 h and showed a maximal response to the addn. of 10-50~mg/l of 2,4-D at 48 h. The promotive effect of 2,4-D was inhibited by cycloheximide suggesting that de novo protein synthesis was involved in this phenomenon. Interestingly, similar promotive effects on the UFGT activity were obsd.

for other phytohormones such as kinetin and several anti-auxins.

94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies

882-09-7, PCIB

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(promotive effect of auxins on UDP-glucose: flavonol

glucosyltransferase activity in Vitis sp. cell cultures)

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

RN 882-09-7 HCAPLUS
CN Propanoic acid, 2-(4-chlorophenoxy)-2-methyl- (9CI) (CA INDEX NAME)

CC

ST plant suspension culture auxin flavonol glucosyltransferase induction

IT Glucosylation (biol.; promotive effect of auxins on UDP-glucose: flavonol glucosyltransferase activity in Vitis sp. cell cultures)

IT Carrot Catharanthus roseus

Cinnamomum cassia
Datura inoxia
Gardenia jasminoides
Grape
Ocimum basilicum
Parsley (Petroselinum crispum)
Patchouli
Rhubarb (Rheum palmatum)
Safflower (Carthamus tinctorius)
Saffron (Crocus sativus)
Stevia rebaudiana
Sweet potato
Tobacco

11-2 (Plant Biochemistry)

(promotive effect of auxins on UDP-glucose: flavonol glucosyltransferase activity in Vitis sp. cell cultures)

IT Hormones, plant
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(promotive effect of auxins on UDP-glucose: flavonol
alucosyltransferase activity in Vitis sp. cell cultures)

glucosyltransferase activity in Vitis sp. cell cultures)
Plant tissue culture

PRYOR 09/781,695

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525-79-1, Kinetin 575-89-3, 2,4,6-T 771-50-6, Indole-3-carboxylic acid
     830-96-6, 1H-Indole-3-propanoic acid 882-09-7, PCIB
     21293-29-8, Abscisic acid
                                 98640-00-7
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (promotive effect of auxins on UDP-glucose: flavonol
        qlucosyltransferase activity in Vitis sp. cell cultures)
     117-39-5, Quercetin
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (promotive effect of auxins on UDP-glucose: flavonol
        glucosyltransferase activity in Vitis sp. cell cultures)
     482-35-9, Quercetin 3-0-glucoside
                                         9075-75-6, UDP-glucose-quercetin
IT
     glucosyltransferase
     RL: BPR (Biological process); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
     nonpreparative); PROC (Process)
        (promotive effect of auxins on UDP-glucose: flavonol
     glucosyltransferase activity in <u>Vitis sp. cell cultures</u>) 6892-74-6, Quercetin 3,7-di-O-glucoside 29125-80-2 1335
                                                29125-80-2 133563-23-2
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
        (promotive effect of auxins on UDP-glucose: flavonol
        glucosyltransferase activity in Vitis sp. cell cultures)
E COUNT: 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L95 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                          2001:597745 HCAPLUS
                          135:148598
DOCUMENT NUMBER:
TITLE:
                          Inducing production of
                          isoflavones in plants using nuclear receptor
                          ligands
INVENTOR(S):
                          Graham, Terrence L.; Graham, Lian-mei Y.; Landini,
                          Serena
                          Ohio State University Research Foundation, USA
PATENT ASSIGNEE(S):
SOURCE:
                          PCT Int. Appl., 46 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                       KIND DATE
                                             APPLICATION NO.
                                                              DATE
                                             WO 2001-US4420
                                                              20010212
     WO 2001058262
                             20010816
                        A1
             AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
             HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
             LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ,
                                          TM, TR, TT,
                                                      TZ, UA, UG, US, UZ, VN,
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                             US 2001-781695
                                                              20010212
     US 2002004458
                             20020110
                        A1
                                          US 2000-181707P P 20000211
PRIORITY APPLN. INFO.:
OTHER SOURCE(S):
                          MARPAT 135:148598
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AB Methods for increasing the levels of isoflavones in plants are provided. The method comprise applying a biol. effective amt. of compn. comprising a select nuclear receptor ligand to the plant. Compns. for inducing the prodn. of isoflavones in plants are also provided. Such compns. comprise one or preferably, a combination of the select nuclear receptor ligands. The nuclear receptor ligand is: (a) a steroid I (rings A and B have the same or different degrees of satn.; R1 = 0 or OR; R2,R5 = H, or Me; R3 = H, OH or O; R4 = R3, CO2H, COCH2OH or COMe; R6 = H, OH or OMe) or II; (b) a phenolic estrogen or di-Ph 4-HOC6H4R7C6H4OH-4 (R7 = bond, alkane or alkene); (c) a long-chain fatty acid R8CO2R9 (R8 = C5-25 aliph. chain; R9 = H or C 1-5 aliph. chain); (d) a peroxisome proliferator R1OR11 R11R12CO2R13 (R10 = arom. ring; R11 = 0 or S; R12, R13 = C1-8 aliph. chain); or (e) zearalenone. The compns. also comprise a compd. that enhances the capacity of the plant to release daidzein and/or utilize it for the prodn. of glyceollin. The action of such a compd. is complementary to that of the nuclear receptor ligand.

II

882-09-7, Clofibric acid 17413-79-5,
2-(2-Chlorophenoxy)-2-methylpropionic acid
RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
(nuclear receptor ligand inducer of isoflavone
prodn. in plants)

RN 882-09-7 HCAPLUS

CN Propanoic acid, 2-(4-chlorophenoxy)-2-methyl- (9CI) (CA INDEX NAME)

RN 17413-79-5 HCAPLUS

CN Propanoic acid, 2-(2-chlorophenoxy)-2-methyl- (9CI) (CA INDEX NAME)

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Me
            - CO2H
         Me
IC
     ICM A01N031-00
     ICS A01N035-00; A01N037-00; A01N037-44; A01N039-02
     5-3 (Agrochemical Bioregulators)
     Section cross-reference(s): 11
ST
     isoflavone prodn plant nuclear receptor ligands
IT
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
         (agonists; nuclear receptor ligand inducer of
        isoflavone prodn. in plants)
IT
     Nuclear receptors
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
         (ligands; inducers of isoflavone prodn.
        in plants)
IT
     Fatty acids, biological studies
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
         (long-chain; nuclear receptor ligand inducer of
        isoflavone prodn. in plants)
TT
     Peroxisome proliferators
        (nuclear receptor ligand inducer of isoflavone
        prodn. in plants)
     Alfalfa (Medicago sativa)
TT
     Bean (Phaseolus limensis)
     Chickpea (Cicer arietinum)
     Peanut (Arachis hypogaea)
     Plant (Embryophyta)
     Soybean (Glycine max)
        (nuclear receptor ligand inducers of isoflavone
        prodn. in)
     Bean (Phaseolus vulgaris)
        (pinto; nuclear receptor ligand inducers of
        isoflavone prodn. in)
     Onium compounds
IT.
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
         (tetrazolium, redox dyes; enhancer of nuclear receptor ligand
        inducers of isoflavone prodn. in plants)
     9012-72-0, glucan
TT
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
(Phytophthora sojae cell wall, fragment; enhancer of nuclear receptor
        ligand inducers of isoflavone prodn. in
        plants)
                                14333-18-7, Orthovanadate
TT
     11121-48-5, Rose bengal
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
         (enhancer of nuclear receptor ligand inducers of
        isoflavone prodn. in plants)
                                50-23-7, Hydrocortisone 50-27-1, Estriol
IT.
     50-02-2, Dexamethasone
     50-28-2, 17.beta.-Estradiol, biological studies 52-39-1, 53-06-5, Cortisone 53-16-7, Estrone, biological studies
                                                           52-39-1, Aldosterone
```

Androsterone 56-53-1, Diethylstilbestrol

57-83-0, Progesterone,

```
prodn. in plants)
     446-72-0D, Genistein, conjugates 485-72-3D, Formononetin, aglycon
     486-66-8D, Daidzein, aglycon 574-12-9, isoflavone
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (nuclear receptor ligand inducers of isoflavone
        prodn. in plants)
L95 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1986:31716 HCAPLUS
DOCUMENT NUMBER:
                         104:31716
TITLE:
                         Formation of chalcones and
                         isoflavones by callus culture of Glycyrrhiza
                         uralensis with different production patterns
                         Kobayashi, Mitsugu; Noguchi, Hiroshi; Sankawa, Ushio
AUTHOR(S):
                         Fac. Pharm. Sci., Univ. Tokyo, Tokyo, 113, Japan
CORPORATE SOURCE:
                         Chemical & Pharmaceutical Bulletin (1985), 33(9),
SOURCE:
                         3811-16
                         CODEN: CPBTAL; ISSN: 0009-2363
DOCUMENT TYPE:
                         Journal
                         English
     Formononetin, isoliquiritigenin, echinatin, liquiritigenin,
     p-hydroxybenzoic acid, 3'-hydroxyformononetin and isobavachalcone were
     isolated from callus culture of Glycyrrhiza uralensis which was
     established on Murashige-Skoog's medium contg. NAA (2 ppm), 2,4-D (1 ppm)
     and benzyladenine (0.1 ppm). Formononetin, 3'-hydroxyformononetin and
     isobavachalcone showed different patterns of prodn.
     94-75-7, biological studies
    RL: BIOL (Biological study)
        (formation of chalcones and isoflavones by callus
        culture of Glycyrrhiza uralensis response to)
     94-75-7 HCAPLUS
    Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
           0- CH2- CO2H
    11-1 (Plant Biochemistry)
     Glycyrrhiza culture chalcone isoflavone; auxin Glycyrrhiza culture
     flavonoid; cytokinin Glycyrrhiza culture flavonoid
IT
    Plant tissue culture
        (formation of chalcone and isoflavones by callus,
        of Glycyrrhiza uralensis, auxins and cytokinins effect on)
IT
    Plant hormones and regulators
     RL: BIOL (Biological study)
        (formation of chalcones and isoflavones by callus
        culture of Glycyrrhiza uralensis response to)
    Ketones, biological studies
TT
     RL: FORM (Formation, nonpreparative)
        (chalcones, formation of, by callus culture of Glycyrrhiza uralensis,
        auxins and cytokinins effect on)
    Flavones
TT
     RL: FORM (Formation, nonpreparative)
        (iso-, formation of, by callus culture of Glycyrrhiza
        uralensis, auxins and cytokinins effect on)
IT
     Licorice
        (G. uralensis, chalcones and is flavones formati n
        by callus culture of, auxins and cytokinins effect on)
     86-87-3 94-75-7, biological studies
                                          1214-39-7
     RL: BIOL (Biological study)
```

PRYOR 09/781,695

(formation of chalcones and isoflav nes by callus culture of Glycyrrhiza uralensis response to)

IT 99-96-7, biological studies 485-72-3 578-86-9 961-29-5 20575-57-9 20784-50-3 34221-41-5

RL: FORM (Formation, nonpreparative) (formation of, by callus culture of Glycyrrhiza uralensis, auxins and cytokinins effect on)

packet #1

PRYOR 09/781,695

=> file reg
FILE 'REGISTRY' ENTERED AT 15:45:35 ON 22 AUG 2003
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STRUCTURE FILE UPDATES: 20 AUG 2003 HIGHEST RN 569883-36-9 DICTIONARY FILE UPDATES: 20 AUG 2003 HIGHEST RN 569883-36-9

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2003

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Experimental and calculated property data are now available. See HELP
     PROPERTIES for more information. See STNote 27, Searching Properties
     in the CAS Registry File, for complete details:
                                                                carno cyclic my
     http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf
     => d que stat 142
                       SCR 1838 AND 2005 AND 1199
     L17
     119
                      SCR 1839 OR 2043
              1018656 SEA FILE=REGISTRY ABB=ON PLU=ON C/RELF AND NRS=1 AND O>1 AND
     L35
                      O<5 AND C<25 AND N<3 NOT (PMS/CI OR (P OR SI)/ELS)
                                                     ~ no pglymers
aromatic
                                                    S @12
                                          0 @7
                 0
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@9 10
        95
     VAR G1=0/12
     VAR G2=7/9
     NODE ATTRIBUTES:
     CONNECT IS X3 RC AT CONNECT IS E2 RC AT
     CONNECT IS E1 RC AT
     CONNECT IS E1 RC AT
     CONNECT IS E2 RC AT
                            12
     DEFAULT MLEVEL IS ATOM
     GGCAT IS UNS AT
     DEFAULT ECLEVEL IS LIMITED
     ECOUNT IS M6 C AT 1
ECOUNT IS X8 C AT 3
ECOUNT IS X5 C AT 10
     GRAPH ATTRIBUTES:
     RING(S) ARE ISOLATED OR EMBEDDED
     NUMBER OF NODES IS 10
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STEREO ATTRIBUTES: NONE

L41 8846 SEA FILE=REGISTRY SUB=L35 SSS FUL L36 AND L17 NOT L19
L42 8472 SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM 8472 cp L5

=> file hcaplus
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This file contains CAS Registry Numbers for easy and accurate substance identification.

search for daim 1

=> d c	que nos 19	(T = 1 m talla
1.17		SCR 1838 AND 2005 AND 1199 SCR 1839 OR 2043 SEA FILE-RECISTRY ARR-ON PLU-ON C/RELE AND NRS-1 AND ON AND termine
L19		SCR 1839 OR 2043
L35	1018656	JEA TILL-REGISTRY ADD-ON TEG-ON C/REET AND MRS-I AND ONL AND
		O<5 AND C<25 AND N<3 NOT (PMS/CI OR (P OR SI)/ELS)
L36		STR
L41		
L42		SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM SEA FILE=HCAPLUS ABB=ON PLU=ON L42 SEA FILE=HCAPLUS ABB=ON PLU=ON "GROWTH AND DEVELOPMENT, NT= narry
L43		SEA FILE=HCAPLUS ABB=ON PLU=ON L42
L46	61035	SEA FILE=HCAPLUS ABB=ON PLU=ON "GROWTH AND DEVELOPMENT, NT= narn
		PLANT"+PFT,NT/CT
L51	3295	SEA FILE=HCAPLUS ABB=ON PLU=ON PEROXISOME PROLIFERATOR-ACTIVA
		TED RECEPTORS+PFT/CT
L52	715	SEA FILE=HCAPLUS ABB=ON PLU=ON PEROXISOME PROLIFERATORS+PFT,N
		т/ст
L53		SEA FILE=HCAPLUS ABB=ON PLU=ON NUCLEAR RECEPTORS+PFT,NT/CT
L63	8175	SEA FILE=HCAPLUS ABB=ON PLU=ON ?FLAVON?(5A)(PRODUC? OR
		INDUC? OR BIOSYNTH? OR FORM?)
L65	27305	SEA FILE=HCAPLUS ABB=ON PLU=ON "PLANT HORMONES AND REGULATORS
		"+PFT,NT/CT
L66		SEA FILE=HCAPLUS ABB=ON PLU=ON L43 AND L63
L67		SEA FILE=HCAPLUS ABB=ON PLU=ON L66 AND (L51 OR L52 OR L53)
L68		SEA FILE=HCAPLUS ABB=ON PLU=ON L66 AND (L46 OR L65)
L70		SEA FILE=HCAPLUS ABB=ON PLU=ON L68 NOT AGEING/TI
L71	1	SEA FILE=HCAPLUS ABB=ON PLU=ON L67 NOT RAT/TI SEA FILE=HCAPLUS ABB=ON PLU=ON 11-5/SC,SX = plant biochemis try
L76	15016	SEA FILE=HCAPLUS ABB=ON PLU=ON 11-5/SC, SX Prant 11-5/SC, SX
L77		SEA FILE=HCAPLUS ABB=ON PLU=ON L43 AND L76
L78		SEA FILE=HCAPLUS ABB=ON PLU=ON L77 AND (DISEAS? OR RESIST?)
L79	1	SEA FILE=HCAPLUS ABB=ON PLU=ON L78 AND (?FLAVON? OR AGLYCON?
		OR GLYCEOLLIN?)
L93		SEA FILE=HCAPLUS ABB=ON PLU=ON (L70 OR L71) OR L79
L94	6	SEA FILE=HCAPLUS ABB=ON PLU=ON L93 AND (CROP OR STEM OR ROOT
		or leaf or foliage) 6 cites

=> d ibib abs hitstr 194 1

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L94 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN
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ACCESSION NUMBER: DOCUMENT NUMBER:

1992:252407 HCAPLUS

116:252407

TITLE:

The biosynthetical capacity of the active principles of "in vitro" regenerated Solenostema argel (Sel.)

Hayne, callus and shoots

AUTHOR(S):

Amariei, Doina; Stanescu, Ursula; Gille, Elvira;

Onisei, Tatiana

CORPORATE SOURCE:

"Stejarul" Res. Stn., Piatra Neamt, 5600, Rom.

SOURCE:

Revue Roumaine de Biologie, Serie de Biologie Vegetale

(1991), 36(1-2), 71-6

CODEN: RRBVD5; ISSN: 0250-5517

DOCUMENT TYPE: Journal LANGUAGE: English

AB The nontoxic active pharmaceutical components of S. argel were previously shown to have anti-inflammatory, anti-ulcerous, and immunostimulatory activity. This report describes the induction of callus and shoot regeneration from S. argel explants in response to various concns. of benzylaminopurine, NAA, and 2,4-D. The active pharmacol. compds. identified in regenerated callus and shoots included flavones, polyphenols, carotenoids, phytosterols, and polyholosides. Their levels were similar to or superior to those in control leaves.

IT 94-75-7, biological studies RL: BIOL (Biological study)

(in induction of Solenostema argel callus and shoot regenerants, biosynthesis of pharmacol.-active compds. in relation to)

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

=> d ind 194 1

L94 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

CC 11-8 (Plant Biochemistry)

ST Solenostema callus shoot regeneration explant hormone; pharmaceutical Solenostema callus regenerated shoot

IT Solenostemma argel

(callus tissue and in vitro regenerated shoots of, hormonal induction of and biosynthesis of pharmacol.-active compds. by)

IT Pharmaceuticals

(formation of, by Solenostema argel callus tissue and in vitro regenerated shoots)

IT Plant hormones and regulators

RL: BIOL (Biological study)

(induction of Solenostema argel callus and shoot regeneration by, biosynthesis of pharmacol.-active compds. subsequent to)

IT Regeneration, biological

(of Solenostema argel shoots, from cultured explants, hormonal induction of)

IT Carotenes and Carotenoids, biological studies

RL: BIOL (Biological study)

(pharmacol.-active, formation of, by Solenostema argel callus and regenerated shoots)

IT Plant tissue

(callus, of Solenostema argel, hormonal induction of and biosynthesis of pharmacol.-active compds. by)

IT Steroids, biological studies

RL: BIOL (Biological study)

(hydroxy, pharmacol.-active, formation of, by Solenostema argel callus and regenerated shoots)

IT Flavonoids

RL: BIOL (Biological study)

(oxo, pharmacol.-active, **formation** of, by Solenostema argel callus and regenerated shoots)

IT Phenols, biological studies

RL: BIOL (Biological study)

(polyhydric, pharmacol.-active, formation of, by Solenostema argel callus and regenerated shoots)

IT Plant tissue

(shoot, Solenostema argel in vitro regenerated, hormonal induction of and biosynthesis of pharmacol. active compds. by) 1214-39-7 TT

86-87-3, NAA **94-75-7**, biological studies

RL: BIOL (Biological study)

(in induction of Solenostema argel callus and shoot regenerants, biosynthesis of pharmacol.-active compds. in relation to)

=> d ibib abs hitstr ind 194 2-6

L94 ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1992:18445 HCAPLUS

DOCUMENT NUMBER:

116:18445

TITLE:

Effects of culture conditions on isoflavonoid levels of transformed and non-transformed cultures of Lupinus

- a comparison of suspension and hairy root

cultures

AUTHOR(S):

Berlin, J.; Ruegenhagen, C.; Rippert, M.; Erdogan, S. Biol. Bundesanst. Forst- und Landwirtsch.,

CORPORATE SOURCE:

Braunschweig, D-3300, Germany

SOURCE:

Zeitschrift fuer Naturforschung, C: Journal of

Biosciences (1991), 46(9-10), 735-42

CODEN: ZNCBDA; ISSN: 0341-0382

DOCUMENT TYPE:

Journal

English LANGUAGE:

Some highly productive suspension and hairy root cultures were found among several transformed cultures of L. polyphyllus and L. hartwegii. A transformed suspension culture Lupo 30150 and a root culture Luha 15834 contg. the highest specific isoflavone glucoside content were characterized and compared with normal phytohormone-dependent lines with respect to product stability as well as their responsiveness to external triggers, e.g. response to changes in the medium. While phytohormone-dependent suspension cultures lost their initial ability to form increased levels of isoflavonoids on phytohormone-free medium, the transformed phytohormone-independent suspension Lupo 30150 remained a highly productive line, despite the fact that its specific levels decreased to 60% of the initial values during several years in liq. medium. Highest stability of product patterns and levels were noted for the transformed root culture. Phytohormones had little effect on growth and isoflavonoid levels in suspension cultures, while they reduced both strongly in root cultures. In the presence of 2,4-D the root culture changed into an aggregated low producing suspension culture from which the root state recovered on phytohormone-free medium. As long as the root state was maintained, isoflavonoid levels could not be distinctly improved by media variation while specific isoflavonoid levels of suspensions were increased by stress factors such as phosphate depletion. When suspensions were transferred to fresh medium phenylalanine ammonia-lyase was greatly induced within 24 h, while the activity remained nearly unchanged in root cultures. IT

94-75-7, biological studies RL: BIOL (Biological study)

(isoflavonoid glycosides formation in lupine

cultures response to)

RN 94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN

11-2 (Plant Biochemistry)

```
isoflavonoid glycoside lysine culture genetic transformation
       Plant hormones and regulators
      .RL: BIOL (Biological study)
          (isoflavonoid glycoside formation and growth of
          lupine tissue cultures response to)
      Plant tissue culture
  IT
          (isoflavonoid glycosides formation by lupine hairy
          roots in)
       Agrobacterium rhizogenes
  IT
       Agrobacterium tumefaciens
          (lupine transformation by, isoflavonoid glycoside
          formation and culture conditions in relation to)
       Transformation, genetic
  IT
          (of lupine, isoflavonoid glycosides formation in
          culture in relation to)
 IT
       Glycosides
       RL: FORM (Formation, nonpreparative)
          (isoflavonoid, formation of, by lupine transformed
          and nontransformed suspension and hairy root cultures)
      <del>-Plant tissue culture</del>
          (suspension, isoflavonoid glycosides formation by
          transformed and nontransformed lupine in)
  IT
          (L. hartwegii, isoflavonoid glycoside formation in
          transformed and nontransformed suspension and hairy root
          cultures of, culture conditions effect on)
  IT
      Lupine
          (L. polyphyllus, isoflavonoid glycoside formation
          in transformed and nontransformed suspension and hairy root
          cultures of, culture conditions effect on)
       14265-44-2, Phosphate, biological studies
. IT
       RL: BIOL (Biological study)
          (deficiency of, isoflavonoid glycoside formation
          and growth of lupine cultures response to)
                    137351-12-3
  IT
       36190-98-4
                                  138110-87-9
       RL: FORM (Formation, nonpreparative)
          (formation of, in lupine cultures, culture conditions effect on)
       57-50-1, Sucrose, biological studies
                                              6484-52-2, Ammonium nitrate,
  IT
       biological studies
                           7757-79-1, Potassium nitrate, biological studies
       RL: BIOL (Biological study)
          (isoflavonoid glycoside formation and growth of
          lupine cultures response to)
       86-87-3, 1-NAA 94-75-7, biological studies
  IT
       1214-39-7
       RL: BIOL (Biological study)
          (isoflavonoid glycosides formation in lupine
          cultures response to)
  L94 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN
  ACCESSION NUMBER:
                           1990:213986 HCAPLUS
  DOCUMENT NUMBER:
                           112:213986
                           Application of growth substances and mineral nutrition
  TITLE:
                           affecting disease development and
                           glyceollin production of soybean
                           Chakraborty, U.; Chakraborty, B. N.; Purkayastha, R.
  AUTHOR(S):
                           Cent. Life Sci., Univ. North Bengal, Darjeeling, 734
  CORPORATE SOURCE:
                           430, India
                           Folia Microbiologica (Prague, Czech Republic) (1989),
  SOURCE:
                           34(6), 490-7
                           CODEN: FOMIAZ; ISSN: 0015-5632
  DOCUMENT TYPE:
                           Journal
  LANGUAGE:
                           English
       The effects of foliar application of growth substances and mineral
       nutrition of the host on the development of charcoal rot disease
       in soybean caused by Macrophomina phaseolina was tested. Among the eight
       growth substances examd., gibberellic acid was most successful in reducing
```



the disease severity, followed by IAA and 2,3,5-triiodobenzoic acid. Low concns. of these compds. stimulated (and high concns. inhibited) the mycelial growth of M. phaseolina in vitro. Substrate supplementation with different doses of N, P, K and Ca had varying effects on disease development. Disease was increased considerably by both excess and deficient N and also by deficient Ca, while excess Ca conferred partial resistance. Glyceollin contents of host roots before and after excess Ca and gibberellic acid (10 mg/L) treatments were estd. significantly increased glyceollin prodn. in infected roots. However, gibberellic acid induced glyceollin synthesis even in uninoculated roots. Changes in the host reaction towards increased resistance was correlated with increased phytoalexin prodn. 94-75-7, 2,4-D, biological studies RL: BIOL (Biological study) (charcoal rot disease of soybean inhibition by) ·94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

IT

RN

CN

CC 11-5 (Plant Biochemistry)

ST soybean infection Macrophomina phytohormone nutrient

IT Plant nutrition

(mineral, charcoal rot disease of soybean development response to growth substances and)

IT Macrophomina phaseolina

(soybean infection by, growth substances and mineral nutrition effect on)

IT Mineral elements

RL: BIOL (Biological study)

(Macrophomina phaseolina growth response to, charcoal rot disease of soybean development in relation to)

IT Soybean

(disease, charcoal rot, phytohormones and mineral nutrition effect on)

IT 77-06-5, Gibberellic acid 87-51-4, IAA, biological studies TIBA 94-75-7, 2,4-D, biological studies

RL: BIOL (Biological study)

(charcoal rot disease of soybean inhibition by)

IT 86-87-3, NAA 120-23-0, 2-Naphthoxyacetic acid 525-79-1, Kinetin 1214-39-7, 6-Benzylaminopurine RL: BAC (Biological activity or effector, except adverse); BSU (Biological

study, unclassified); BIOL (Biological study) (charcoal rot disease of soybean response to)

IT 57103-57-8D, derivs.

RL: FORM (Formation, nonpreparative)

(formation of, by soybean in charcoal rot disease, growth substances and mineral nutrition effect on)

TT 7440-09-7, Potassium, biological studies 7440-70-2, Calcium, biological studies 7723-14-0, Phosphorus, biological studies 7727-37-9, Nitrogen, biological studies

RL: BIOL (Biological study)

(Macrophomina phaseolina growth response to nutrient, charcoal rot disease of soybean in relation to)

L94 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1988:628692 HCAPLUS

DOCUMENT NUMBER:

109:228692

PRYOR 09/781,695 TITLE: Initiation and maintenance of callus tissue culture of Uncaria elliptica for flavonoid producti n Law, K. H.; Das, N. P. AUTHOR(S): Fac. Med., Natl. Univ. Singapore, Singapore, 0511, CORPORATE SOURCE: Progress in Clinical and Biological Research (1988), SOURCE: 280(Plant Flavonoids Biol. Med. 2: Biochem., Cell., Med. Prop.), 67-70 CODEN: PCBRD2; ISSN: 0361-7742 DOCUMENT TYPE: Journal LANGUAGE: Enalish Using specific combinations of auxin and cytokinin hormones, calli initiation and growth were found to occur on young leaf explants of the Uncaria elliptica plant. There was also a different pattern of flavonoid prodn. found in the calli tissue. The major flavonoid, rutin, was not found in the calli tissue but was present in the source plant. However, (-)-epicatechin was detected in almost equal amts. both in the calli tissue and the source plant. The formation of this flavonoid occurred when the calli were grown in the dark. An increase in kinetin concn. showed a moderate increase in epicatechin accumulation. 94-75-7, Dichlorophenoxyacetic acid, biological studies RL: BIOL (Biological study) (flavonoid prodn. enhancement by, in callus tissue culture of Uncaria elliptica) 94-75-7 HCAPLUS RN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN 0-CH2-CO2H 16-6 (Fermentation and Bioindustrial Chemistry) CC Uncaria callus tissue culture flavonoid auxin; cell culture Uncaria epicatechin prodn cytokinin IT Uncaria elliptica (callus tissue culture of, flavonoid prodn. with, plant hormones effect on) ΙT Flavonoids RL: BMF (Bioindustrial manufacture); BIOL (Biological study); PREP. (Preparation) (manuf. of, by callus tissue culture of Uncaria elliptica, auxins and cytokinins effect on) Plant hormones and regulators RL: BIOL (Biological study) (auxins, flavonoid prodn. enhancement by, in callus tissue culture of Uncaria elliptica)

IT Plant tissue culture (callus, flavonoid prodn. by, of Uncaria elliptica, auxins and cytokinins effect on) IT Plant hormones and regulators RL: BIOL (Biological study) (cytokinins, flavonoid prodn. enhancement by, in callus tissue culture of Uncaria elliptica) 86-87-3, .alpha.-Naphthalene acetic acid 94-75-7, Dichlorophenoxyacetic acid, biological studies 133-32-4 Indole-3-butyric acid 525-79-1, Kinetin (plant hormone) 133-32-4, 1214-39-7, N6-Benzyladenine 2365-40-4 RL: BIOL (Biological study) (flavon id prodn. enhancement by, in callus tissue culture of Uncaria elliptica)

490-46-0P, (-)-Epicatechin RL: BMF (Bioindustrial manufacture); BIOL (Biological study); PREP (Preparation) (manuf. of, by callus tissue culture of Uncaria elliptica, auxins and cytokinins effect on) L94 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 1986:512153 HCAPLUS DOCUMENT NUMBER: 105:112153 Effects of 5% maltose and plant growth regulators on TITLE: the callus growth and flavonoid formation of some Scutellaria baicalensis stem callus lines Yamamoto, Hisako; Chatani, Nobuyasu; Watanabe, Kume; AUTHOR(S): Tomimori, Tsuyoshi Sch. Pharm., Hokuriku Univ., Kanazawa, 920-11, Japan CORPORATE SOURCE: Shoyakugaku Zasshi (1986), 40(1), 33-9 SOURCE: CODEN: SHZAAY; ISSN: 0037-4377 DOCUMENT TYPE: Journal LANGUAGE: Japanese Studies were conducted on effects of plant growth regulators on the growth and flavonoid content of some S. baicalensis stem callus lines on Linsmaier-Skoog medium contg. 5% maltose instead of 3% sucrose. optimum combination of plant growth regulators in line St-20 were 10-7M NAA and 10-5M kinetin, and 10-5M NAA and 10-5M kinetin for callus growth and 10-6M IAA and 10-5M kinetin, and 10-7M NAA and 10-5M kinetin for flavonoid content. When 5% maltose was added to the medium, an increase in flavonoid content was obsd. in line St-20 but not in 3 other lines. 94-75-7, biological studies IT RL: BIOL (Biological study) (callus growth and flavonoid formation response to, in Scutellaria baicalensis) 94-75-7 HCAPLUS RN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN 0-CH2-CO2H 11-3 (Plant Biochemistry) Scutellaria callus growth flavonoid phytoregulator; maltose callus growth ST Scutellaria IT Plant hormones and regulators RL: BIOL (Biological study) (callus growth and flavonoid formation response to, in Scutellaria baicalensis) IT Plant tissue culture (callus, flavonoid formation and growth of, of Scutellaria baicalensis, phytoregulators effect on) IT Scutellaria baicalensis (flavonoid formation and callus growth of, phytoregulators effect on) Flavonoids IT RL: FORM (Formation, nonpreparative) (formation of, by Scutellaria baicalensis callus cultures, phytoregulators effect on) IT Glycosides RL: FORM (Formation, nonpreparative) (flavonoid, formation of, by Scutellaria

baicalensis callus cultures, phytoregulators effect on) 50-99-7, biological studies 57-50-1, biological studies

87-51-4, biological studies 94-75-7, biological

IT

86-87-3

69-79-4

studies 525-79-1 RL: BIOL (Biological study) (callus growth and flav n id formati n response to, in Scutellaria baicalensis) 21967-41-9 480-40-0 491-67-8 632-85-9 36948-76-2 51059-44-0 57396-78-8 104125-36-2 RL: FORM (Formation, nonpreparative) (formation of, in Scutellaria baicalensis callus culture, carbon sources and phytoregulators effect on) L94 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 1986:203955 HCAPLUS DOCUMENT NUMBER: 104:203955 TITLE: Flavonoid production in Scutellaria baicalensis callus cultures AUTHOR(S): Yamamoto, Hisako; Chatani, Nobuyasu; Kitayama, Akemi; Tomimori, Tsuyoshi CORPORATE SOURCE: Sch. Pharm., Hokuriku Univ. Kanagawa, Kanazawa, 920-11, Japan SOURCE: Plant Cell, Tissue and Organ Culture (1986), 5(3), 219-22 CODEN: PTCEDJ; ISSN: 0167-6857 DOCUMENT TYPE: Journal LANGUAGE: English St-20 and St-7 lines were isolated from the stem callus of S. baicalensis Georgi on indole-3-acetic acid and 2,4-dichlorophenoxyacetic acid media, resp. The flavonoid content of St-20 line was superior to that of St-7 line. The growth and flavonoid (baicalin, baicalein, wogonin, wogonin-7-0-glucuronide) contents in St-20 line were best on Linsmaier-Skoog's basal medium contg. 10-7M-10-5M kinetin. St-20 line showed the same flavonoid content and pattern as the root of parent plant after 70 days of culturing. 94-75-7, biological studies RL: BIOL (Biological study) (growth and flavonoid formation by Scutellaria baicalensis callus response to) 94-75-7 HCAPLUS Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN 0-CH2-CO2H 11-2 (Plant Biochemistry) CC Section cross-reference(s): 16 Scutellaria callus flavonoid ST Plant growth and development (by Scutellaria baicalensis callus, flavonoid formation in relation to) IT Plant tissue culture (callus, of Scutellaria baicalensis, flavanoid formation in, phytohormones effect on) TT Plant hormones and regulators RL: BIOL (Biological study) (flavonoid formation by callus cultures of Scutellaria baicalensis response to) Scutellaria baicalensis (flavonoid formation by callus of, phytohormones effect on) IT Flavonoids

RL: FORM (Formation, nonpreparative)

(formation of, by Scutellaria baicalensis callus,

PRYOR 09/781,695

phytohormones effect on)

IT 491-67-8 632-85-9 21967-41-9 51059-44-0

RL: FORM (Formation, nonpreparative)
 (formation of, by Scutellaria baicalensis callus culture, phytohormones effect on)

IT 87-51-4, biological studies 94-75-7, biological studies 525-79-1

RL: BIOL (Biological study)
 (growth and flavonoid formation by Scutellaria baicalensis callus response to)

=> s 193 not 194
L95 6 L93 NOT L94 4— eites cover elaim => 1 but

=> d ibib abs hitstr ind 15 1-6 are directed to

L5 ANSWER 1 OF 12 HCAPLUS COPYRIGHT 2003 ACS on STN plant cultures

See parket # 2 for

Here references

PRYOR 09/781,695

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=> d que nos 183
                SCR 1838 AND 2005 AND 1199
L17
                SCR 1839 OR 2043
L19
        1018656 SEA FILE=REGISTRY ABB=ON PLU=ON C/RELF AND NRS=1 AND O>1 AND
L35
                O<5 AND C<25 AND N<3 NOT (PMS/CI OR (P OR SI)/ELS)
L36
                STR
           8846 SEA FILE=REGISTRY SUB=L35 SSS FUL L36 AND L17 NOT L19
141
L42
           8472 SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM
          32167 SEA FILE=HCAPLUS ABB=ON PLU=ON L42
L43
L76
          15016 SEA FILE=HCAPLUS ABB=ON PLU=ON 11-5/SC,SX
             60 SEA FILE=HCAPLUS ABB=ON PLU=ON L43 AND L76
L77
             23 SEA FILE=HCAPLUS ABB=ON PLU=ON L77 AND (DISEAS? OR RESIST?)
L78
              2 SEA FILE=HCAPLUS ABB=ON PLU=ON L78 AND (COMBINATION OR
L83
                COMPOSITION)
                                      2 cites
=> d que nos 184
                SCR 1838 AND 2005 AND 1199
L17
<del>L19</del>
                SCR 1839 OR 2043
        1018656 SEA FILE=REGISTRY ABB=ON PLU=ON C/RELF AND NRS=1 AND O>1 AND
L35
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L36
L41
           8846 SEA FILE=REGISTRY SUB=L35 SSS FUL L36 AND L17 NOT L19
L42
           8472 SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM
L43
          32167 SEA FILE=HCAPLUS ABB=ON PLU=ON L42
            215 SEA FILE=HCAPLUS ABB=ON PLU=ON L43 AND (?FLAVON? OR AGLYCON?
L74
                OR GLYCEOLLIN?)
              7 SEA FILE=HCAPLUS ABB=ON PLU=ON L74 AND DISEAS?
L75
L76
          15016 SEA FILE=HCAPLUS ABB=ON PLU=ON 11-5/SC,SX
              1 SEA FILE=HCAPLUS ABB=ON PLU=ON L75 AND L76 1 cite
L84
=> d que nos 187
                SCR 1838 AND 2005 AND 1199
L17
L19
                SCR 1839 OR 2043
        1018656 SEA FILE=REGISTRY ABB=ON PLU=ON C/RELF AND NRS=1 AND O>1 AND
L35
                O<5 AND C<25 AND N<3 NOT (PMS/CI OR (P OR SI)/ELS)
L36
                STR
1.41
           8846 SEA FILE=REGISTRY SUB=L35 SSS FUL L36 AND L17 NOT L19
           8472 SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM
L42
          32167 SEA FILE=HCAPLUS ABB=ON PLU=ON L42
143
            215 SEA FILE=HCAPLUS ABB=ON PLU=ON L43 AND (?FLAVON? OR AGLYCON?
L74
                OR GLYCEOLLIN?)
             19 SEA FILE=HCAPLUS ABB=ON PLU=ON L74 AND (DISEAS? OR RESIST?)
L85
              7 SEA FILE=HCAPLUS ABB=ON PLU=ON L85 AND (PLANT OR CROP OR
L86
                STEM OR ROOT OR LEAF OR FOLIAGE)
              6 SEA FILE=HCAPLUS ABB=ON PLU=ON L86 NOT AGEING/TI 6 cites
L87
=> d que nos 192
                SCR 1838 AND 2005 AND 1199
L17
                SCR 1839 OR 2043
L19
        1018656 SEA FILE=REGISTRY ABB=ON PLU=ON C/RELF AND NRS=1 AND O>1 AND
L35
                O<5 AND C<25 AND N<3 NOT (PMS/CI OR (P OR SI)/ELS)
L36
                STR
           8846 SEA FILE=REGISTRY SUB=L35 SSS FUL L36 AND L17 NOT L19
L41
           8472 SEA FILE=REGISTRY ABB=ON PLU=ON L41/COM
L42
          32167 SEA FILE=HCAPLUS ABB=ON PLU=ON L42
L43
        1995008 SEA FILE=HCAPLUS ABB=ON PLU=ON IRON OR ORTHOVANDATE OR ROSE
L88
                BENGAL OR TETRAZOLIUM OR COPPER OR CU OR FE OR PHYTOALEXIN OR
                GLUCAN OR PHYTOPHTHORA
L89
           1130 SEA FILE=HCAPLUS ABB=ON PLU=ON L88 AND L43
             50 SEA FILE=HCAPLUS ABB=ON PLU=ON L89 AND DISEAS?
L90
             24 SEA FILE=HCAPLUS ABB=ON PLU=ON L90 AND (PLANT OR CROP OR
L91
                STEM OR ROOT OR LEAF OR FOLIAGE)
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L92

21 SEA FILE=HCAPLUS ABB=ON PLU=ON L91 NOT (PHARMACEUTICAL OR 21 cites workers or liver)/TI

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=> s 183-84 or 187 or 192
L96 27 (L83 OR L84) OR L87 OR L92 27 cites total
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=> d ibib abs hitstr ind 1-27

L96 ANSWER 1 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

2003:13172 HCAPLUS

DOCUMENT NUMBER:

138:133384

TITLE:

Somatic embryo formation and plant

regeneration in 'Zaoh' line No.2 of Japanese angelica

tree (Aralia elata seem.)

AUTHOR(S):

Amemiya, Keiichi; Mochizuki, Tohru

CORPORATE SOURCE:

Yamanashi Agricultural Research Center, Yamanashi,

405-0105, Japan

SOURCE:

Plant Biotechnology (Tokyo, Japan) (2002), 19(5),

383-387

303-301

CODEN: PLBIF6; ISSN: 1342-4580

PUBLISHER: DOCUMENT TYPE: Japanese Society for Plant Cell and Molecular Biology

DOCUMENT TYPE: Journal LANGUAGE: English

AB Mass propagation of the Japanese angelica tree (Aralia elata seem.) 'Zaoh' line No.2 was established through somatic embryos. Petioles of leaflets were cultured for induction of calli on an MS medium contg. 1 mgL-1 of

2,4-D in combination with 0.5 mgL-1 BA. The initiated calli were moved onto MS medium supplemented with glutamine 450 mgL-1 and asparagine 300 mgL-1. An embryogenic callus developed after 5 mo and was moved onto regulator-free MS medium. Numerous plantlets were regenerated from this embryogenetic callus, and rooted plantlets were potted after acclimation and planted to the field. The plants had the same characteristics in morphol. in the field, except for the no. of thorns per

internode. The plants also had the same resistance to Phytophthora disease as was in the original genotype

'Zaoh' line No.2. This study provided a method for mass propagation from petioles of leaflets and proves that regenerated plants maintain the morphogenetic characteristics and disease resistance of the original genotype.

T 94-75-7, 2,4-D, biological studies

RL: BSU (Biological study, unclassified); BIOL (Biological study)
(MS medium contg.; somatic embryo formation and plant
regeneration in 'Zaoh' line No.2 of Japanese angelica tree (Aralia elata seem.))

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

CC 9-11 (Biochemical Methods)
 Section cross-reference(s): 11

ST Aralia somatic embryo formation plant regeneration

IT Plant tissue culture

(callus; somatic embryo formation and plant regeneration in 'Zaoh' line No.2 of Japanese angelica tree (Aralia elata seem.))

IT Aralia elata Regeneration, plant

Somatic embryogenesis, plant

(somatic embryo formation and plant regeneration in 'Zaoh'

line No.2 of Japanese angelica tree (Aralia elata seem.)) IT 94-75-7, 2,4-D, biological studies 1214-39-7, BA RL: BSU (Biological study, unclassified); BIOL (Biological study) (MS medium contg.; somatic embryo formation and plant regeneration in 'Zaoh' line No.2 of Japanese angelica tree (Aralia elata seem.)) 70-47-3, Asparagine, biological IT 56-85-9, Glutamine, biological studies studies RL: BSU (Biological study, unclassified); BIOL (Biological study) (MS medium supplemented with; somatic embryo formation and plant regeneration in 'Zaoh' line No.2 of Japanese angelica tree (Aralia elata seem.)) L96 ANSWER 2 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 2001:8659 HCAPLUS DOCUMENT NUMBER: 134:190774 TITLE: Arabidopsis dth9 mutation identifies a gene involved in regulating disease susceptibility without affecting salicylic acid-dependent responses AUTHOR(S): Mayda, Esther; Mauch-Mani, Brigitte; Vera, Pablo Instituto de Biologia Molecular y Celular de Plantas, CORPORATE SOURCE: Universidad Politecnica-Consejo Superior de Investigaciones Cientificas, Valencia, 46022, Spain SOURCE: Plant Cell (2000), 12(11), 2119-2128 CODEN: PLCEEW; ISSN: 1040-4651 PUBLISHER: American Society of Plant Physiologists DOCUMENT TYPE: Journal LANGUAGE: English To det. which components of the plant defense response make important contributions to limiting pathogen attack, an M2 mutagenized population of a transgenic Arabidopsis line was screened for mutants showing constitutive expression of .beta.-glucuronidase activity driven by the promoter region of the CEVI-1 gene. The CEVI-1 gene originally was isolated from tomato plants and has been shown to be induced in susceptible varieties of tomato plants by virus infection in a salicylic acid-independent manner. The authors report here the characterization of a recessive mutant, detachment9 (dth9). This mutant is more susceptible to both virulent and avirulent forms of the oomycete Peronospora and also exhibits increased susceptibility to the moderately virulent bacterial pathogen Pseudomonas syringae pv maculicola ES4326. However, this mutant is not affected in salicylic acid metab. and shows normal expression of pathogenesis-related (PR) genes after pathogen attack. Furthermore, after inoculation with avirulent pathogens, the dth9 mutant shows a compromised systemic acquired resistance response that cannot be complemented by exogenous application of salicylic acid, although this mol. is able to promote normal activation of PR genes. Therefore, the dth9 mutation defines a regulator of disease susceptibility that operates upstream or independently of salicylic acid. Pleiotropy is also evident in the dth9 mutant in the sense that the shoots of dth9 plants are insensitive to the exogenously applied auxin analog 2,4-dichlorophenoxyacetic acid. 94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(Arabidopsis with mutation in gene involved in regulating disease susceptibility without affecting salicylic acid-dependent responses insensitivity to)

RN 94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

```
0-CH2-CO2H
     11-5 (Plant Biochemistry)
     Section cross-reference(s): 3
     Arabidopsis mutation gene disease susceptibility salicylate
ST
     dependence
IT
     Chromosome
        (Arabidopsis thaliana 2; of DTH9 gene involved in regulating
        disease susceptibility without affecting salicylic
        acid-dependent responses linkage in Arabidopsis)
IT
     Phytoalexins
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
        (Arabidopsis with mutation in gene involved in regulating
        disease susceptibility without affecting salicylic
        acid-dependent responses accumulation of)
IT
     Auxins
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (Arabidopsis with mutation in gene involved in regulating
        disease susceptibility without affecting salicylic
        acid-dependent responses insensitivity to)
IT
     Gene, plant
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (DTH9; mutation in gene involved in regulating disease
        susceptibility without affecting salicylic acid-dependent responses of
        Arabidopsis)
IT
     Mutation
        (in gene involved in regulating disease susceptibility
        without affecting salicylic acid-dependent responses of Arabidopsis)
IT
     Arabidopsis thaliana
        (mutation in gene involved in regulating disease
        susceptibility without affecting salicylic acid-dependent responses of)
TT
     Disease resistance, plant
     Peronospora parasitica
     Pseudomonas syringae maculicola
        (mutation in gene involved in regulating disease
        susceptibility without affecting salicylic acid-dependent responses of
        Arabidopsis)
IT
        (of DTH9 gene involved in regulating disease susceptibility
        without affecting salicylic acid-dependent responses in Arabidopsis)
IT
     135531-86-1, Camalexin
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
        (Arabidopsis with mutation in gene involved in regulating
        disease susceptibility without affecting salicylic
        acid-dependent responses accumulation of phytoalexin)
TT
     94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (Arabidopsis with mutation in gene involved in regulating
        disease susceptibility without affecting salicylic
        acid-dependent responses insensitivity to)
     69-72-7, Salicylic acid, biological studies
RL: BAC (Biological activity or effector, except adverse); BSU (Biological
```

study, unclassified); BIOL (Biological study)

(mutation in gene involved in regulating disease

susceptibility without affecting salicylic acid-dependent responses of

Arabidopsis)

REFERENCE COUNT:

THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS 33 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L96 ANSWER 3 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1999:587780 HCAPLUS

DOCUMENT NUMBER:

132:47704

TITLE:

Induced resistance and phytoalexin

accumulation in biological control of early blight

disease for tomato plants by gamma irradiation and growth regulators

AUTHOR(S):

El-Sayed, S. A.; El-Hawa, Abou

CORPORATE SOURCE:

Radiobiology Department, Nuclear Research Center,

Atomic Energy Commission, Egypt

SOURCE:

Pakistan Journal of Biochemistry and Molecular Biology

(1996), 29(1-2), 41-50

CODEN: PJBBF5

PUBLISHER:

Pakistan Society of Biochemistry and Molecular Biology

DOCUMENT TYPE:

lournal LANGUAGE: Enalish

Gibberellic acid (GA), indole 3-acetic acid (IAA), 6-furfurylaminopurine (kinetin), 2,4-dichlorophenoxyacetic acid (2,4-D), and Alar were found to be abiotic elicitors for a phytoalexin, rishitin, in tomato and represented a crit. role in its accumulation. On the contrary, gamma irradn. was not able to initiate rishitin formation. Application with GA and IAA, followed by artificial infection with Alternaria solani, had a synergistic effect on rishitin accumulation, which led to a highly significant resistance to early blight caused by A. solani. Meanwhile, application with growth inhibitors, 2,4-D and Alar, as well as gamma irradn. treatment followed by infection with A. solani had an antagonistic effect on rishitin accumulation that led to significant increase in susceptibility to early blight. Hence, it was concluded that pre-infectional application with GA and/or LAA could induce and/or improve early blight disease resistance in tomato. The 2,4-D, Alar and/or gamma irradn. tend to weaken natural immunity by lowering the biogenerating capacity of tomato leaves and the accumulation of the natural antibiotic, rishitin.

94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(induced resistance and phytoalexin accumulation in biol.

control of early blight disease for tomato plants

by gamma irradn. and growth regulators)

94-75-7 HCAPLUS RN

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN

11-5 (Plant Biochemistry) CC

Section cross-reference(s): 10

blight tomato rishitin phytohormone gamma ray ST

(early blight; induced resistance and phytoalexin accumulation in biol. control of early blight disease for tomato plants by gamma irradn. and growth regulators)

IT Alternaria solani

Gamma ray

Tomato

(induced resistance and phytoalexin accumulation in biol. control of early blight disease for tomato plants

```
by gamma irradn. and growth regulators)
IT
     Hormones, plant
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (induced resistance and phyt alexin accumulation in biol.
        control of early blight disease for tomato plants
        by gamma irradn. and growth regulators)
IT
     Phytoalexins
     RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological
     study, unclassified); BIOL (Biological study); OCCU (Occurrence); PROC
        (induced resistance and phytoalexin accumulation in biol.
        control of early blight disease for tomato plants
        by gamma irradn. and growth regulators)
     77-06-5, Gibberellic acid 87-51-4, Indole 3-Acetic acid, biological
     studies 94-75-7, 2,4-Dichlorophenoxyacetic acid, biological
               525-79-1, 6-Furfurylaminopurine 1596-84-5, Alar
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (induced resistance and phytoalexin accumulation in biol.
        control of early blight disease for tomato plants
        by gamma irradn. and growth regulators)
     18178-54-6, Rishitin
IT
     RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological
     study, unclassified); BIOL (Biological study); OCCU (Occurrence); PROC
     (Process)
        (induced resistance and phytoalexin accumulation in biol.
        control of early blight disease for tomato plants
        by gamma irradn. and growth regulators)
                               THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         54
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L96 ANSWER 4 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1997:106411 HCAPLUS
DOCUMENT NUMBER:
                          126:140630
TITLE:
                         Significance and application of microbial toxicity
                         tests in assessing ecotoxicological risks of
                         contaminants in soil and sediment
                         van Beelen, P.; Doelman, P.
AUTHOR(S):
                         National Inst. Public Health Environment, Bilthoven,
CORPORATE SOURCE:
                         3720 BA, Neth.
SOURCE:
                         Chemosphere (1997), 34(3), 455-499
                         CODEN: CMSHAF; ISSN: 0045-6535
PUBLISHER:
                         Elsevier
                         Journal
DOCUMENT TYPE:
LANGUAGE:
                         .English
    Micro-organisms are vital for soil fertility and for the degrdn. of org.
     matter and pollutants in soils and sediments. Due to their function and
     ubiquitous presence micro-organisms can act as an environmentally very
     relevant indicator of pollution. Microbial tests should be used
     discriminatory for the establishment of soil and sediment quality
     guidelines. This review gives an evaluation of microbial toxicity tests
     and a novel method to derive quality guidelines. Long term microbial
     tests are generally less sensitive than short term tests.
     effects can be obscured by the activity of a few resistant
     micro-organisms, when for example soil respiration is used as a sum
     parameter during a long incubation period. Mineralization tests with high
     substrate concns. which enable growth, are less sensitive than similar tests with low concns. of substrate. The latter tests are more relevant
```

for natural ecosystems. The often applied microbial toxicity tests can be categorized as single species tests, biomass measurements, carbon and nitrogen transformations, enzymic tests and tests measuring changes in microbial diversity. Comparisons between tests can only be indicative because the relative sensitivity depends on the toxicants and soils used. The respiration rate per unit of biomass is a more sensitive indicator of toxic effects than the respiration rate or the amt. of biomass alone. The autotrophic nitrification and acetylene redn. tests can be sensitive when

short incubation times are used. The nitrogen mineralization, denitrification and many enzymic tests are often not very sensitive. The urease activity is a relatively sensitive enzymic test in many studies. The replacement of sensitive micro-organisms by different resistant species can have serious ecol. consequences. Some species become extinct while others appear in bulging nos. Adaptation of a community to a pollutant must be considered as the very process which disturbs a polluted ecosystem. The resistant micro-organisms often fail to perform specific ecol. functions. The occurrence of resistant species can be used as an sensitive and ecol. relevant indicator for deterioration from environmental pollution. Persistent toxic effects on the microflora can be caused by zinc, cadmium and copper at concn. levels lower than European Community limits. Tests with anaerobic sediment processes were orders of magnitude more sensitive for some chlorinated aliph. compds. than aquatic toxicity tests. The addn. of a few mg zinc per kg soil can inhibit the more sensitive microbial processes (like chloroform or 4-chlorophenol degrdn.), whereas soil invertebrates and some plants are less sensitive to zinc. After the evaluation of the tests, a novel method is described to derive soil and sediment quality guidelines using microbial toxicity tests. The results of single species tests with micro-organisms can be incorporated into the contemporary risk assessment method for higher organisms which is based on the extrapolation from single species tests to the protection of 95% of all species in an ecosystem. This method uses the No Obsd. Effect Concns. (NOEC) of a no. of toxicity tests to calc. a Hazardous Concn. 5% (HC5). The HC5 is calcd. from more than 5 NOEC values. In analogy the Effect Concn. 10% (EC10) can be used to calc. the Dangerous Concn. 5% (DC5). The DC5 is calcd. from more than 5 EC10 values. The DC5 should give protection to 95% of the microbial processes. The DC5 of a no. of pollutants are calcd. and compared with the HC5 values from the literature. Microbial toxicity tests can be used for risk assessment because micro-organisms are among the most sensitive organisms for the effects of pollutants.

94-75-7, biological studies

RL: ADV (Adverse effect, including toxicity); POL (Pollutant); BIOL

(Biological study); OCCU (Occurrence)

(application of microbial toxicity tests in assessing ecotoxicol, risks of contaminants in soil and sediment)

94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN

CC 4-1 (Toxicology)

microbial toxicity test ecotoxicity pollutant; soil pollution ecotoxicity microbial toxicity test; sediment contaminant ecotoxicity microbial toxicity test; ecotoxicol risk pollutant test microbial toxicity

IT Aquatic sediments Ecotoxicity **Nitrification** Nitrogen cycle Respiration, microbial Risk assessment Soil pollution Toxicity

Water pollution (application of microbial toxicity tests in assessing ecotoxicol. risks of contaminants in soil and sediment)

IT Trace metals RL: ADV (Adverse effect, including toxicity); POL (Pollutant); BIOL

```
(Biological study); OCCU (Occurrence)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
ΙŢ
     Enzymes, biological studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
    Toxicants
        (contaminants; application of microbial toxicity tests in assessing
        ecotoxicol. risks of contaminants in soil and sediment)
IT
     Trace metals
     RL: ADV (Adverse effect, including toxicity); POL (Pollutant); BIOL
     (Biological study); OCCU (Occurrence)
        (heavy; application of microbial toxicity tests in assessing
        ecotoxicol. risks of contaminants in soil and sediment)
    Bioassay
TT
        (microbial toxicity test; application of microbial toxicity tests in
        assessing ecotoxicol, risks of contaminants in soil and sediment)
IT
    Heavy metals
     RL: ADV (Adverse effect, including toxicity); POL (Pollutant); BIOL
     (Biological study); OCCU (Occurrence)
        (trace; application of microbial toxicity tests in assessing
        ecotoxicol. risks of contaminants in soil and sediment)
     67-66-3, Chloroform, biological studies
                                               71-43-2, Benzene, biological
IT
               25323-30-2, Dichloroethylene
     studies
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
TT
     56-38-2, Parathion
                          62-53-3, Aniline, biological studies
     Benzoic acid, biological studies
                                       69-72-7, 2-Hydroxybenzoic acid,
                         74-82-8, Methane, biological studies 74-8-tudies 74-85-1, Ethene, biological studies
     biological studies
     Ethane, biological studies
     74-99-7, Propyne 76-87-9, Triphenyltin hydroxide
                                                           87-86-5
                        93-07-2, 3,4-Dimethoxybenzoic acid 94-75-7,
     Pentachlorophenol
                         95-47-6, Orthoxylene, biological studies
     biological studies
     3,4-Dichloroaniline
                          99-05-8, 3-Aminobenzoic acid
                                                          99-06-9,
     3-Hydroxybenzoic acid, biological studies 99-10-5, 3,5-Dihydroxybenzoic
            100-09-4, 4-Methoxybenzoic acid
                                              100-71-0, 2-Ethylpyridine
     101-21-3, Chlorpropham 106-47-8, 4-Chloroaniline, biological studies
     107-00-6, 1-Butyne 108-42-9, 3-Chloroaniline 118-92-3, 2-Aminacid 149-91-7, 3,4,5-Trihydroxybenzoic acid, biological studies
                                                      118-92-3, 2-Aminobenzoic
     150-13-0, 4-Aminobenzoic acid
                                     150-68-5
                                                490-79-9, 2,5-Dihydroxybenzoic
                                           536-74-3, Phenylacetylene
            501-65-5, Diphenyl acetylene
                          584-13-4, 4-Amino-1,2,4-triazole
     542-75-6, Telone II
                                                              625-53-6
                    627-19-0, 1-Pentyne 1420-07-1, Dinoterb
     Ethylthiourea
                                                                 1423-60-5
                     1504-58-1, 3-Phenyl-2-propyn-1-ol
                                                        1918-16-7, Propachlor
     3-Butyn-2-one
     1929-82-4, Nitrapyrin 1945-84-2, 2-Ethynylpyridine
                                                            2921-88-2,
     Chlorpyrifos
                   4187-87-5, 1-Phenyl-2-propyn-1-ol 4685-14-7, Paraquat
     7439-92-1, Lead, biological studies
                                           7439-96-5, Manganese, biological
     studies
               7440-02-0, Nickel, biological studies
                                                       7440-38-2, Arsenic,
                          7440-43-9, Cadmium, biological studies
    biological studies
     Chromium, biological studies 7440-50-8, Copper, biological studies
     7440-66-6, Zinc, biological studies
                                           12427-38-2, Maneb
                                                                12789-03-6,
                 13194-48-4, Ethoprop
     Chlordane
                                       14998-27-7, Chlorite
                                                                15972-60-8.
                             16520-62-0, 4-Phenyl-1-butyne
                                                              19044-88-3,
                16088-73-6
     Alachlor
                21609-90-5, Leptophos
                                       21725-46-2, Cyanazine
     Oryzalin
                       50594-66-6, Acifluorfen
                                                 51218-45-2, Metolachlor
     Methylpyrimifos
     51338-27-3
                  53780-34-0, Mefluidide
                                            55283-68-6, Ethalfluralin
                                            69806-40-2, Haloxyfop methyl
     58138-08-2, Tridiphane
                              66546-20-1
                  81412-43-3, Tridemorph
     69806-50-4
                                           81777-89-1, Dimethazone
     87818-31-3, Cinmethylin
     RL: ADV (Adverse effect, including toxicity); POL (Pollutant); BIOL
     (Biological study); OCCU (Occurrence)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
```

59-67-6, Nicotinic acid, biological studies

IT

63-91-2, L-Phenylalanine,

PRYOR 09/781,695

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103-82-2, Phenylacetic acid, biological studies
                 120-80-9, 1,2-Benzenediol, biological studies
     Ouercetin
                                                                      138-52-3
                156-38-7, 4-Hydroxyphenylacetic acid Flavone 592-57-4, 1,3-Cyclohexadiene
     Salicine
                                                           495-69-2, Benzoylglycine
     525-82-6, Flavone
                                                           7400-08-0.
     4-Hydroxycinnamic acid
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
     56-40-6, Glycine, biological studies
                                               56-65-5, biological studies
     56-86-0, Glutamic acid, biological studies
                                                    64-19-7, Acetic acid,
     biological studies
                           9002-13-5, Urease 9016-17-5, Arylsulfatase
     37341-58-5, Phytase
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (application of microbial toxicity tests in assessing ecotoxicol. risks
        of contaminants in soil and sediment)
     74-86-2, Acetylene, biological studies
                                                 <del>7439-89-6, Iron, biological</del>
     studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (redn.; application of microbial toxicity tests in assessing
        ecotoxicol. risks of contaminants in soil and sediment)
L96 ANSWER 5 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                           1992:632985 HCAPLUS
DOCUMENT NUMBER:
                           117:232985
TITLE:
                           Rare earth metals-containing compositions for growth
                           stimulation and disease prevention, in
                           plants.
                           Ning, Jiagong; Li, Guangming; Liu, Sui; et al.
INVENTOR(S):
                           Hunan Research Center of Rare Earth Agricultural
PATENT ASSIGNEE(S):
                           Application, Peop. Rep. China
SOURCE:
                           Faming Zhuanli Shenqing Gongkai Shuomingshu, 16 pp.
                           CODEN: CNXXEV
DOCUMENT TYPE:
                           Patent
LANGUAGE:
                           Chinese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                       KIND
                              DATE
                                              APPLICATION NO.
                                                                DATE
     CN 1061888
                        Α
                              19920617
                                              CN 1990-106134
                                                                19901208
     CN 1034273
                        В
                              19970319
PRIORITY APPLN. INFO.:
                                           CN 1990-106134
                                                                19901208
     The title compns. consists of rare earth compds., trace elements,
     plant growth regulators, buffers, surfactants, and
     membrane-forming agents. A compn. for rice consisted of Ce salt 0-40, La
     salt 0-40, La salt-Y salt mixt. 0-40, Zn and Zn salt 5-40, boric acid 5-20, Fe salt 0-20, Mn salt 0-40, carboxylig-acid 5-20, starch
     0-10, surfactant 0-15, growth regulator 0-1, and 2,4-D 0-1 g. Compared to
     conventional Ce salts, these formulations produced 30-80% higher yield.
     94-75-7, 2,4-D, biological studies
     RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study);
     USES (Uses)
        (growth promotion and disease prevention by compns. contg.,
        in plants)
     94-75-7 HCAPLUS
     Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
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biological studies 100-21-0, 1,4-Benzenedicarboxylic acid, biological

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0-CH2-CO2H
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ICM A01N059-20

19-6 (Fertilizers, Soils, and Plant Nutrition)

Section cross-reference(s): 5

rare earth plant growth stimulator; disease protection plant rare earth

Plant hormones and regulators RL: BIOL (Biological study)

(growth promotion and disease prevention by compns. contg.

rare earth metals and, in plants)

Carboxylic acids, biological studies Rare earth metals, biological studies Trace elements, biological studies

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(growth promotion and disease prevention by compns. contg.,

in plants) Plant disease

IT (prevention of, by compns. contg. rare earth metals) 77-06-5 94-75-7, 2,4-D, biological studies 557-34-6, Zinc IT

1344-67-8, Copper chloride 7429-90-5, Aluminum, 7439-89-6, Iron, biological studies biological studies 7439-89-6D, **Iron**, salts 7439-91-0D, Lanthanum, salts

7439-95-4, Magnesium, biological studies 7439-96-5D, Manganese, salts 7439-98-7, Molybdenum, biological studies 7440-00-8D, Neodymium, salts 7440-32-6, Titanium, biological studies 7440-10-0D, Praseodymium, salts 7440-42-8, Boron, biological studies 7440-45-1D, Cerium, salts

7440-50-8, Copper, biological studies 7440-65-5D, Yttrium,

7440-66-6, Zinc, biological studies 7440-66-6D, Zinc, salts salts 7487-88-9, Magnesium sulfate, biological studies 7646-85-7, Zinc

chloride, biological studies 7733-02-0, Zinc sulfate 7758-94 Ferrous chloride 7758-98-7, Copper sulfate, biological studies 7733-02-0, Zinc sulfate 7758-94-3,

7785-87-7, Manganese sulfate 7786-30-3, 7773-01-5, Manganese chloride Magnesium chloride, biological studies 9005-25-8, Starch, biological

10402-29-6, **Copper** nitrate 11098-84-3, Ammonium studies

14013-86-6, Ferrous nitrate molvbdate

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(growth promotion and disease prevention by compns. contg., in plants)

L96 ANSWER 6 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

CORPORATE SOURCE:

1992:588364 HCAPLUS

DOCUMENT NUMBER:

117:188364

TITLE:

Comparison of plant hormone requirements in leaf

tissues from hop stunt viroid-infected and uninfected

hop plants

AUTHOR(S):

SOURCE:

Takahashi, T.; Fujiwara, S.; Chiba, K.; Yoshikawa, N.

Fac. Agric., Iwate Univ., Morioka, 020, Japan

Zeitschrift fuer Pflanzenkrankheiten und Pflanzenschutz (1992), 99(1), 62-70

CODEN: ZPFPAA; ISSN: 0340-8159

DOCUMENT TYPE:

Journal

LANGUAGE: German

The effects of various auxin/cytokinin ratios on the growth of leaf segments from hop stunt viroid-infected hop plants and healthy controls were studied in culture expts. Thus, c mbinations of

indole-3-acetic acid (I) with 6-benzylaminopurine (II) promoted tissue growth in both infected and uninfected leaves; for the former the effect was greater, however. Kinetin (III) combinati ns with I exhibited a similar, but smaller effect, as did combinations of indole-3-butyric acid and II or III and naphthalene-1-acetic acid with II or III. 2.4-Dichlorophenoxyacetic acid combinations with II or III resulted in almost as large a growth stimulation as I with II or III. Of I precursors tested with II, indole was particularly effective in promoting growth. Gibberellic acid in combination with II or I+II promoted growth particularly in noninfected leaves. Rooting was depressed in infected leaf cultures, even under I+II combinations promoting leaf tissue growth. Effects of viroid infection on auxin metab., esp. on I are discussed. 94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies RL: BIOL (Biological study) (in hop leaf growth and rooting, hop stunt viroid infection effect on) 94-75-7 HCAPLUS Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

CC 11-5 (Plant Biochemistry)
Section cross-reference(s): 10

ST viroid infection hop growth auxin kinetin; hormone plant viroid infection hop growth

IT Leaf

Root

RN CN

(growth of, in hop, hop stunt virus infection inhibition of, auxins and other hormones in)

IT Plant hormones and regulators RL: BIOL (Biological study)

(in hop leaf growth and rooting, hop stunt viroid infection effect on)

IT Plant growth and development
(leaf growth, in hop, hop stunt virus infection inhibition of, auxins and other hormones in)

IT Plant hormones and regulators
RL: BIOL (Biological study)

(auxins, in \hat{h} op leaf growth and rooting, hop stunt viroid infection effect on)

IT Viroid

(hop stunt, hop plant infection with, leaf growth and rooting inhibition by, auxins and other hormones in)

IT Plant growth and development

(rooting, in hop, hop stunt virus infection inhibition of, auxins and other hormones in)

IT Hop

(H. lupulus, disease, hop stunt virus infection, leaf growth and rooting inhibition by, auxins and other hormones in)

L96 ANSWER 7 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1992:510792 HCAPLUS

DOCUMENT NUMBER:

117:110792

TITLE:

Potato yields and nitrate accumulation

AUTHOR(S):

Abazov, A. Kh.; Urtaev, D. A.

PRYOR 09/781,695

CORPORATE SOURCE:

NPO Kartofelevod., USSR

SOURCE:

Khimizatsiya Sel'skogo Khozyaistva (1988-1992) (1991),

(7), 17-19

CODEN: KSKHE7; ISSN: 0235-2516

DOCUMENT TYPE:

Journal

LANGUAGE:

Russian

Spraying a \underline{DTPA} complex of Fe + Cu + Zn + Co during flower bud formation, followed by spraying 10 g 2M-4Kh/ha in 30 kg/ha of a N 10-P 34 kg/ha urea-contg. liq. fertilizer 20-25 days before harvest or Mg chlorate 10-12 days before harvest, increased potato yield and decreased tuber NO3-. 2M-4Kh and Mg chlorate were ineffective under a late blight infection.

TT

94-74-6, MCPA RL: BIOL (Biological study)

(potato tuber nitrate control by haulm desiccation by)

RN 94-74-6 HCAPLUS

Acetic acid, (4-chloro-2-methylphenoxy)- (9CI) (CA INDEX NAME)

0- CH2-CO2H

19-5 (Fertilizers, Soils, and Plant Nutrition) CC

Section cross-reference(s): 5, 17

potato nitrate trace metal chlorate MCPA ST

Food contamination IT

(by nitrates, of potatoes, trace metal fertilization and haulm desiccation control of)

Potato IT

(fertilizer expt. with, with nitrate contamination control by trace metals and haulm desiccation)

Plant desiccants IT

(potato tuber nitrate control by)

TT Phytophthora

(potato tuber nitrate response to haulm desiccation and infection by)

Fertilizer experiment IT

(with trace metals, with potato, tuber nitrate in relation to)

TT Potato

> (disease, late blight, tuber nitrate response to haulm desiccation and)

IT Trace elements, compounds

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses) (metals, complexes, fertilizer expt. with, with potato, tuber nitrate in relation to)

67-43-6D, DTPA, trace metal complexes TT

15162-64-8 65229-17-6 142198-25-2 RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)

(fertilizer expt. with, with potato, tuber nitrate control in relation to)-

IT 14797-55-8, Nitrate, biological studies

RL: BIOL (Biological study)

(in potatoes, trace metal fertilization and haulm desiccation control

94-74-6, MCPA 10326-21-3, Magnesium chlorate IT

RL: BIOL (Biological study)

(potato tuber nitrate control by haulm desiccation by)

ACCESSION NUMBER:

L96 ANSWER 8 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN 1992:146049 HCAPLUS

116:146049

DOCUMENT NUMBER: TITLE:

Effect of mineral element, hormone and urea on stripe

disease infection in barley AUTHOR(S): Mathur, A. K.; Bhatnagar, G. C. Agric. Res. Stn., Dep. Plant Pathol., Jaipur, 302 018, CORPORATE SOURCE: India Indian Journal of Mycology and Plant Pathology (1990), SOURCE: 20(2), 192-3 CODEN: IJMPAK; ISSN: 0303-4097 Journal DOCUMENT TYPE: LANGUAGE: English 2.4-D. and urea were evaluated against stripe of barley. All Minerals. treatments (except Mo) were consistently superior to the control in reducing disease incidence. In and B were most effective, followed by Fe and Cu. The 2,4-D and urea treatments were less effective. IT 94-75-7, 2,4-D, biological studies RL: BIOL (Biological study) (stripe disease of barley response to, yield in relation to) RN 94-75-7 HCAPLUS Acetic acid. (2.4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN 0- CH2- CO2H 5-2 (Agrochemical Bioregulators) Section cross-reference(s): 19

barley stripe disease mineral element dichlorophenoxyacetate; chlorophenoxyacetate barley stripe disease; urea barley stripe disease IT Mineral elements Plant hormones and regulators RL: BIOL (Biological study) (stripe disease of barley response to, yield in relation to) IT Barley (disease, stripe, control of, mineral elements and hormones and urea in relation to) 57-13-6, Urea, biological studies 94-75-7, 2,4-D, biological IT 7487-88-9, Magnesium sulfate, biological studies 7733-02-0, Zinc sulfate 7758-98-7, Copper Ferrous sulfate sulfate, biological studies 7785-87-7, Manganese sulfate

7720-78-7, 10043-35-3, Boric acid, biological studies 11098-84-3, Ammonium molybdate RL: BIOL (Biological study) (stripe disease of barley response to, yield in relation to)

L96 ANSWER 9 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1990:213986 HCAPLUS

DOCUMENT NUMBER:

112:213986

TITLE:

Application of growth substances and mineral nutrition

affecting disease development and glyceollin production of soybean

AUTHOR(S):

Chakraborty, U.; Chakraborty, B. N.; Purkayastha, R.

CORPORATE SOURCE:

Cent. Life Sci., Univ. North Bengal, Darjeeling, 734

430, India

SOURCE:

Folia Microbiologica (Prague, Czech Republic) (1989),

34(6), 490-7

CODEN: FOMIAZ; ISSN: 0015-5632

DOCUMENT TYPE:

Journal

LANGUAGE:

English

The effects of foliar application of growth substances and mineral AB nutrition of the host on the development of charcoal rot disease in soybean caused by Macrophomina phaseolina was tested. Among the eight

growth substances examd., gibberellic acid was most successful in reducing the disease severity, followed by TAA and 2,3,5-triiodobenzoic acid. Low concns. of these compds. Stimulated (and high concns. inhibited) the mycelial growth of M. phaseolina in vitro. Substrate supplementation with different doses of N, P, K and Ca had varying effects on disease development. Disease was increased considerably by both excess and deficient N and also by deficient Ca, while excess Ca conferred partial resistance. Glyceollin contents of host roots before and after excess Ca and gibberellic acid (10 mg/L) treatments were estd. Both significantly increased glyceollin prodn. in infected roots. However, gibberellic acid induced glyceollin synthesis even in uninoculated roots. Changes in the host reaction towards increased resistance was correlated with increased phytoalexin prodn.

94-75-7, 2,4-D, biological studies

RL: BIOL (Biological study)

(charcoal rot disease of soybean inhibition by)

94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

CC 11-5 (Plant Biochemistry)

soybean infection Macrophomina phytohormone nutrient

TT Plant nutrition

(mineral, charcoal rot disease of soybean development response to growth substances and)

Macrophomina phaseolina IT

(soybean infection by, growth substances and mineral nutrition effect on)

Mineral elements

RL: BIOL (Biological study)

(Macrophomina phaseolina growth response to, charcoal rot disease of soybean development in relation to)

IT Soybean

(disease, charcoal rot, phytohormones and mineral nutrition effect on)

77-06-5, Gibberellic acid 87-51-4, IAA, biological studies TIBA 94-75-7, 2,4-D, biological studies RL: BIOL (Biological study)

(charcoal rot disease of soybean inhibition by)

120-23-0, 2-Naphthoxyacetic acid 86-87-3, NAA 525-79-1, Kinetin 1214-39-7, 6-Benzylaminopurine

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(charcoal rot disease of soybean response to) 57103-57-8D, derivs.

IT

RL: FORM (Formation, nonpreparative)

(formation of, by soybean in charcoal rot disease, growth substances and mineral nutrition effect on)

, Potassium, biological studies 7440-70-2, Calcium, biological 7723-14-0, Phosphorus, biological studies 7727-37-9, Nitrogen, ΙT 7440-09-7, Potassium, biological studies studies biological studies

RL: BIOL (Biological study)

(Macrophomina phaseolina growth response to nutrient, charcoal rot disease of soybean in relation to)

L96 ANSWER 10 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 1990:134371 HCAPLUS

DOCUMENT NUMBER:

112:134371

TITLE:

Effect of various fungicides on mycelial growth, sporangial production, enzyme activity and control of

Phytophth ra leaf blight of Colocasia esculenta L

AUTHOR(S):

Aggarwal, A.; Mehrotra, R. S.

CORPORATE SOURCE:

Bot. Dep., Kurukshetra Univ., Kurukshetra, 132 119,

India

SOURCE:

Acta Phytopathologica et Entomologica Hungarica

(1988), 23(3-4), 401-14

CODEN: APEHEG; ISSN: 0238-1249

DOCUMENT TYPE:

LANGUAGE:

Journal English

AB In studies on the effect of 23 fungicides on mycelial growth of P. colocasiae, Apron 350 FW, Blitox, Blimix, Cuman-L, Demosan 65W, Dexon, Difolatan 80W, Fytolan, Hexaferb, Kitazin, Milton, Ridomil 25 WP and Syllit, showed 100% inhibition at different concns. All the fungicides had some effect on sporangial formation. Studying the effect of 8 fungicides on pectolytic and cellulolytic enzyme activities revealed that Ridomil-25 WP gave max. enzyme inhibition followed by Apron-350FW, Demosan-65W, Difolatan-80W, Phytoalexin 84, Blimix, Fytolan and Topsin-M. Out of 23 systemic and non-systemic fungicides, 8 were tried in

Topsin-M. Out of 23 systemic and non-systemic fungicides, 8 were tried in the field, and Ridomil 25WP at 200 ppm was the most effective, followed by Apron 350FW (500 ppm), Demosan 65W (20 ppm), Difolatan 80W (50 ppm), Phytoalexin-84 (500 ppm), Blimix (100 ppm), Fytolan (200 ppm) and Topsin-M (500 ppm).

IT 2008-39-1, Monosan

RL: BIOL (Biological study)

(Phytophthora colocasiae growth inhibition by)

RN 2008-39-1 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)-, compd. with N-methylmethanamine (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 124-40-3 CMF C2 H7 N

H₃C-NH-CH₃

CM 2

CRN 94-75-7 CMF C8 H6 C12 O3

CC 5-2 (Agrochemical Bioregulators)

ST fungicide Phytophthora Colocasia; sporulation Phytophthora fungicide; taro leaf blight fungicide

IT Phytophthora colocasiae

(control of, on taro, sporulation and enzymes inhibition in)

IT Phytoalexins

RL: BIOL (Biological study)

(taro leaf blight control by, enzyme and sporulation

inhibition in)

IT Fungicides and Fungistats

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(agrochem., taro leaf blight control by, enzyme and
        sporulation inhibition in)
     Colocasia esculenta
IT
         (disease, leaf blight, control of)
ΙT
     Enzymes
     RL: BIOL (Biological study)
         (pectolytic, of Phytophthora colocasiae, fungicide effect on)
     9012-54-8, Cellulase 9015-75-2
IT
                                           9032-75-1, Polygalacturonase
     9033-35-6, Pectin methyl transeliminase 37213-52-8, Poly(Methyl
     galacturonase)
     RL: BIOL (Biological study)
         (of Phytophthora colocasiae, fungicide effect on)
                            2425-06-1, Difolatan 2675-77-6, Demosan
     1332-40-7, Fytolan
īΤ
     23564-05-8
                    56448-75-0, Blimix
                                          57837-19-1, Ridomil
     RL: BIOL (Biological study)
         (taro leaf blight control by, enzyme and sporulation
         inhibition in)
TT
     82-68-8, Brassicol
                                                  137-30-4, Cuman-L
                            133-06-2, Hexacap
                                                                         140-56-7,
     Dexon 2008-39-1, Monosan 2439-10-3, Syllit
Dithane M-45 8066-21-5, Miltox 10605-21-7
                                                         8018-01-7,
                                                         12122-67-7, Dithane Z-78
     13286-32-3, Kitazin
                             14484-64-1, Hexaferb
                                                     17804-35-2. Benlate
     81412-43-3, Calixin
     RL: BIOL (Biological study)
         (Phytophthora colocasiae growth inhibition by)
L96 ANSWER 11 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                           1989:404427 HCAPLUS
DOCUMENT NUMBER:
                           111:4427
TITLE:
                           Low concentrations of phytoalexins correlate
                           with resistance in regenerated
                           plants from meristem cultures of Vicia faba L
AUTHOR(S):
                           Thynn, M.; Wolff, A.; Goerge, E.; Werner, D.
CORPORATE SOURCE:
                           Bot. Inst., Philipps Univ., Marburg, D-3550, Fed. Rep.
SOURCE:
                           Zeitschrift fuer Naturforschung, C: Journal of
                           Biosciences (1989), 44(3-4), 237-42
                           CODEN: ZNCBDA; ISSN: 0341-0382
DOCUMENT TYPE:
                            Journal
LANGUAGE:
                           English
     In tissue cultures from shoot apex meristems with leaf primordias of V. faba, addn. of low concn. of auxins (0.01 mg/L) induced regeneration of whole plants at high frequency (100%). The
     combination of NAA and kinetin/or GA3 also induced a high yield of
     plant regeneration. Regenerated plants from various cultivars on a medium with 2,4-D (0.01 mg/L) were infected with Botrytis
     cinerea, Phytophthora megasperma and Rhizoctonia solani.
     Accumulation of phytoalexins, ethylene prodn. and the
     resistance to fungal diseases were studied. In general,
     prodn. of phytoalexins occurred at a high level in all cultivars
     infected with B. cinerea. Ethylene prodn. varied more in the 7 cultivars
     studied than phytoalexin accumulation. No cultivar was
     resistant to B. cinerea. The highest resistance and the
     lowest concn. of phytoalexin was found after infection by R.
     solani, and phytoalexin accumulation and resistance
     were intermediate in plants infected by P. megasperma. The da
suggest that only low to medium concns. of phytoalexim in faba
     beans are correlated with resistance of regenerated
     plants.
IT
     94-75-7, 2,4-Dichlorophenoxyacetic acid, biological studies
     RL: BIOL (Biological study)
         (regeneration of Vicia faba from meristem cultures in response to)
RN
     94-75-7 HCAPLUS
     Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
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0- CH2- CO2H
     11-5 (Plant Biochemistry)
    Vicia meristem culture auxin regeneration; phytoalexin Vicia
ST
     regnerated plant fungal resistance; Botrytis
     resistance Vicia ethylene phytoalexin; Rhizoctonia
     resistance Vicia regnerated plant phytoalexin;
     Phytophthora resistance Vicia regnerated plant
     phytoalexin
    Plant tissue culture
        (of Vicia faba meristem, phytohormone effects on, phytoalexins
        and resistance to fungal infection of regenerated
       plants in relation to)
IT
     Broad bean
        (phytoalexin accumulation and resistance to fungal
        infection by regenerated plants from meristem cultures of)
IT
     Botrytis cinerea
      Phytophthora megasperma
     Rhizoctonia solani
        (resistance of Vicia faba cultivars regenerated from meristem
        culture to, phytoalexin accumulation in relation to)
     Phytoalexins
     RL: BIOL (Biological study)
        (Vicia faba cultivars regenerated from meristem culture, in response to
        fungal infection, fungal resistance in relation to)
     Plant hormones and regulators
     RL: BIOL (Biological study)
        (auxins, regeneration of Vicia faba meristem cultures in response to)
                           20450-52-6, Wyerol
                                              20450-54-8, Dihydrowyerone
     20079-30-5, Wyerone
TT
     117783-52-5, Wyeronic acid
     RL: FORM (Formation, nonpreparative)
        (formation of, by Vicia faba cultivars regenerated from meristem
        culture, fungal resistance in relation to)
    74-85-1, Ethylene, biological studies
ΙT
     RL: FORM (Formation, nonpreparative)
        (formation of, by Vicia faba cultivars regenerated from meristem
        culture, in response to infection with Botrytis cinerea)
               86-87-3, 1-Naphthaleneacetic acid
                                                  87-51-4, Indole-3-acetic
IT
     acid, biological studies 94-75-7, 2,4-Dichlorophenoxyacetic
                               525-79-1, Kinetin
     acid, biological studies
                                                   1214-39-7.
     6-Benzyl-aminopurine
    RL: BIOL (Biological study)
        (regeneration of Vicia faba from meristem cultures in response to)
L96 ANSWER 12 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1984:98181 HCAPLUS
DOCUMENT NUMBER:
                         100:98181
TITLE:
                         Control of brown spot disease of rice
                         seedlings by treatment with a selected group of
                         chemicals
                         Giri, D. N.; Sinha, A. K.
AUTHOR(S):
                         Dep. Plant Pathol., Bidhan Chandra Krishi
CORPORATE SOURCE:
                         Viswavidyalaya, Kalyani, 741235, India
                         Zeitschrift fuer Pflanzenkrankheiten und
SOURCE:
                         Pflanzenschutz (1983), 90(5), 479-87
                         CODEN: ZPFPAA; ISSN: 0340-8159
DOCUMENT TYPE:
                         Journal
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English

Chems. known to induce the formation of phyt alexins, i.e. Na

malonate [141-95-7], Na molybdate, Na iodoacetate [305-53-3], Na2SO3,

LANGUAGE:

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NaF, as well as DL-methionine [59-51-8] and the phytohormones IAA [87-51-4] and (2,4-D) [94-75-7], reduced brown spot disease symptom expression in-pot-grown (rice) inoculated with Helminthosporium oryzae at 2 days after the treatment. Similar res
                                                                               Similar results
      were shown by root dip at transplanting and by wet seed
      treatment. Leaf diffusates from noninoculated treated seedlings
      had no effect on germ tube growth of H. oryzae. However, diffusates from
      inoculated, treated plants showed germ tube growth inhibition,
      when compared to diffusates from untreated inoculated plants.
      94-75-7, reactions
IT
      RL: RCT (Reactant); RACT (Reactant or reagent)
          (brown spot disease expression redn. by, in rice)
      94-75-7 HCAPLUS
RN
      Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
C1
```

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0-CH2-CO2H
```

5-2 (Agrochemical Bioregulators) Helminthosporium rice phytoalexin induction; malonate rice Helminthosporium; molybdate rice Helminthosporium; iodoacetate rice Helminthosporium **Phytoalexins** IT RL: BIOL (Biological study)

(inducers of, brown spot disease expression response to treatment by, in rice)

IT **Helminthosporium** (rice infection by, phytoalexin-inducers and growth regulators effect on)

IT Rice (disease, brown spot, phytoalexin-inducers and growth regulators effect on)

87-51-4, reactions **94-75-7**, reactions IT RL: RCT (Reactant); RACT (Reactant or reagent) (brown spot disease expression redn. by, in rice)

IT 141-95-7 305-53-3 1313-82-2, reactions 7631-95-0 7681-49-4.

RL: BIOL (Biological study) (phytoalexin induction by, brown spot disease expression redn. by, in rice)

L96 ANSWER 13 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1982:521946 HCAPLUS

DOCUMENT NUMBER:

97:121946

TITLE:

SOURCE:

Metabolism of 2,4-dichlorophenoxyacetic acid in

2,4-dichlorophenoxyacetic acid-resistant

soybean callus tissue

AUTHOR(S):

Davidonis, Gayle H.; Hamilton, Robert H.; Mumma, Ralph

CORPORATE SOURCE:

Pestic. Res. Lab., Pennsylvania State Univ.,

University Park, PA, 16802, USA Plant Physiology (1982), 70(1), 104-7

CODEN: PLPHAY; ISSN: 0032-0889

DOCUMENT TYPE:

English

Journal LANGUAGE:

Three 2,4-D [94-75-7]-resistant root callus tissue lines of Glycine max var Acme were derived by culturing callus tissue 2-6 mo on 40 mg/L 2,4-D and designated 40R, 40B, and 40C. Tissue line 40R had a lower level of 2,4-D uptake in 2-wk-old tissue which disappeared in 3.5-wk-old tissue and less free 2,4-D following incubation for 24 h with [1-14C]2,4-D. This tissue line accumulated more

hydroxylated glycosides of 2,4-D than did nonresistant tissue. Tissue line 40B showed no difference in uptake of 2,4-D when compared to nonresistant tissue, but it did contain less free 2,4-D and more hydroxylated glycosides. The metab. of 2,4-D in the 40C tissue line did not differ significantly from nonresistant tissue, although uptake was less. The 40R line reverted to the same 2,4-D sensitivity as Acme root callus following 6 transfers on 10 .mu.M naphthaleneacetic acid [86-87-3] medium. The altered 2,4-D uptake and metab. characteristic of 40R were also lost. The levels of amino acid conjugates of 2,4-D in the resistant root callus tissue lines were either lower or not significantly different from the Acme tissue lines. Therefore, variations in uptake and metab. of 2,4-D do not wholly explain the resistance of the derived tissue lines, and perhaps modification of the active site or compartmentation is involved. 94-75-7, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (metab. of, in soybean callus culture, herbicidal tolerance in relation

RN 94-75-7 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN

5-3 (Agrochemical Bioregulators) Section cross-reference(s): 11

ST dichlorophenoxyacetate metab soybean callus resistance

IT Soybean

(2,4-D metab. by 2,4-D-resistant callus tissue of)

Plant tissue culture

(2,4-D metab. by soybean callus in, herbicidal tolerance in relation to)

Aglycons IT

RL: BIOL (Biological study)

(dichlorophenoxyacetate metabolites, in soybean callus culture, herbicidal tolerance in relation to)

IT Glycosides

RL: BIOL (Biological study)

(of dichlorophenoxyacetate, in soybean callus culture, herbicidal tolerance in relation to)

IT 3004-84-0

RL: BIOL (Biological study)

(aglycon, of soybean callus tissue, 2,4-D metab. and herbicidal tolerance in relation to)

94-75-7, biological studies

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

-(metab. of, in soybean callus culture, herbicidal tolerance in relation to)

86-87-3 93-76-5

RL: BIOL (Biological study)

(soybean callus culture growth response to, dichlorophenoxyacetate metab. in relation to)

32773-59-4 35144-55-9

RL: BIOL (Biological study)

(soybean callus tissue metabolite, dichlorophenoxyacetate metab. and herbicidal tolerance in relation to)

L96 ANSWER 14 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN 1981:437062 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER:

95:37062

TITLE:

Development of an integrated system to protect apple

orchards from pests and diseases

AUTHOR(S):

Isin, M. M.

CORPORATE SOURCE: SOURCE:

Nauchno-Proizvod. Ob'edin. "Almaly", USSR Vestnik Sel'skokhozyaistvennoi Nauki Kazakhstana

(1981), (4), 44-50

CODEN: VSNKBD; ISSN: 0042-4684

DOCUMENT TYPE:

Journal

LANGUAGE:

Russian

Spraying apples with 0.5% entobacterin 3 plus 0.02% phosalone-(I) [2310-17-0] decreased fruit damage by codling moth by 92.4%. This I dose, 10-fold decreased in comparison with I-alone treatment, 3.5-fold increased the beneficial predator population in comparison with combining the entobacterin-I treatment with 3 releases of Trichogramma, totaling in 5500 wasps/tree, and increased the Trichogramma effectiveness from 4.7 to 6.7%. Adding 1% colloidal S or 0.5% Thiovit [7704-34-9], to the entobacterin-I mixt. controlled powdery mildew and fruit mites, whereas on addn. 0.5% Cu oxycholoride, zineb [12122-67-7], or khomitsin [8066-21-5] controlled apple scab without killing the beneficial insects. Shifting the entobacterin-insecticide sprayings from day- to night decreased the apple damage by codling moth from 12 to 5.6%. Nighttime spraying prevented apple leaf burn by chlorophos [52-68-6]. Replacing soil cultivation by planting perennial grasses decreased the mite infestation from 3.4 to 2.1 mites/leaf, fruit damage by codling moth from 15 to 12.1%, the percentage of cytosporosis-infected trees from 92.8 to 55.2% and the intensity of cytosporosis infection from 52.7 to 23.4%. At 50, 100, and 150 kg N/ha, cytosporosis infection was 12.5, 28.7, and 38.8%, resp., under concomitant application of 50 kg P plus 50 kg K/ha, and even more without the P-K fertilization. K increased the resistance to cytosporosis more than did P. Application of 50 kg N plus 50 kg P/ha, alone or with 5 kg K/ha, decreased the mite infestation from 12.8 to 6.7 and 3.6 mites/leaf, resp. Weed control in apple tree rows with 8 kg simazine [122-34-9] or 1.5 kg 2,4-D amine 2008-39-1]/ha decreased cytosporosis infection intensity from 21.0 to 11.3%. Wintering apple scab was controlled by 10% NH4NO3 or KNO3, or 7% urea. Mite and leaf-eating insect control stimulated apple spur and foliage surface growth and increased fruit vitamin C [50-81-7] and dry matter. Fundazol (II) [17804-35-2] controlled most fungi and the fruit mites. IT

2008-39-1

RL: BIOL (Biological study)

(apple cytosporosis response to weed control by)

2008-39-1 HCAPLUS

Acetic acid, (2,4-dichlorophenoxy)-, compd. with N-methylmethanamine (1:1) CN (9CI) (CA INDEX NAME)

CM

CRN 124-40-3 CMF C2 H7 N

```
CM
          2
     CRN 94-75-7
     CMF C8 H6 C12 O3
           0- CH2- CO2H
CC
     5-4 (Agrochemicals)
     Section cross-reference(s): 19
     apple acaricide insecticide fungicide herbicide
ST
IT
     Genetics
        (apple cytosporosis resistance and)
IT
     Irrigation
     Weed control
        (apple cytosporosis response to)
IT
     Cytospora
        (apple resistance to, increase of)
IT
     Fertilizers
     RL: BIOL (Biological study)
        (apple scab control by)
IT
    Trichogramma
     Entobacterins
     RL: BIOL (Biological study)
        (codling moth control by, on apples, phosalone enhancement of)
IT
     Codling moth
        (control of, biol. and chem.)
IT
     Apple scab
        (control of, by fungicides and fertilizers)
     Erysiphaceae
     Fruit mite
        (control of, on apples)
IT
        (diseases and pests control on, integration of)
IT
     Fungicides and Fungistats
     Insecticides
        (for apples)
IT
    Plant growth and development
        (fundazol and phosalone stimulation of, in apples)
IT
    Ecology
        (in apple disease and pesticide integrated control)
     Apple
IT
        (disease, cytosporosis, resistance to, increase of)
IT
     Fertilizers
     RL: BIOL (Biological study)
        (nitrogen-phosphorus-potassium, apple cytosporosis response to)
     122-34-9 2008-39-1
IT
     RL: BIOL (Biological study)
        (apple cytosporosis response to weed control by)
IT
     1332-40-7
                 8066-21-5
                             12122-67-7
     RL: BIOL (Biological study)
        (apple scab control by)
     7704-34-9, biological studies
                                      17804-35-2
IT
```

RL: BIOL (Biological study)

IT

2310-17-0

(fungi and fruit mites control by, on apples)

H₃C-NH-CH₃

```
RL: BIOL (Biological study)
        (insect and mite control by, on apples)
     52-68-6
    RL: BIOL (Biological study)
        (insect control by, on apples)
     50-81-7, biological studies
     RL: BIOL (Biological study)
        (of apple fruit, pest control by phosalne increase of)
L96 ANSWER 15 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
                         1980:439307 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         93:39307
                         Studies on the foot-rot and leaf-rot
TITLE:
                         diseases of pan (Piper betel Linn.). XI.
                         Effect of plant growth-regulators on
                         endogenous mycelial respiration and growth of
                         Phytophthora parasitica var. Piperina
                         Chaurasia, S. C.
AUTHOR(S):
                         Dep. Bot., Grad. Coll., Seoni, 480661, India
CORPORATE SOURCE:
                         Biochemistry and Experimental Biology (1979), 15(1),
SOURCE:
                         17-24
                         CODEN: BEXBBO; ISSN: 0366-0060
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
    Of 5 plant growth-regulators_tested in .vitro at 1-100 ppm,
     gibberellic acid [77-06-5] caused the highest respiration inhibition of
    P. parasitica piperina mycelium. Indole-3-propionic acid [830-96-6] was
    most inhibitory for radial growth. The fungus was isolated from
     diseased pan.
     94-75-7, biological studies
TT
     RL: BIOL (Biological study)
        (mycelial respiration and growth inhibition by, in Phytophthora
        parasitica piperina)
    94-75-7 HCAPLUS
RN
    Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
           0-CH2-CO2H
     5-2 (Agrochemicals)
     plant growth regulator Phytophthora
TT
    Microorganism respiration
        (by Phytophthora parasitica piperina, plant
        growth-regulators inhibition of)
IT
    Plant hormones and regulators
     RL: BIOL (Biological study)
        (mycelial respiration and growth inhibition by, in Phytophthora
       parasitica piperina)
    Phytophthora parasitica piperina
        (mycelial respiration and growth of, plant growth regulators
        inhibition of)
              87-51-4, biological studies 94-75-7, biological
     77-06-5
              133-32-4
                          830-96-6
     studies
     RL: BIOL (Biological study)
        (mycelial respiration and growth inhibition by, in Phytophthora
        parasitica piperina)
L96 ANSWER 16 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1979:552620 HCAPLUS
```

DOCUMENT NUMBER:

TITLE:

91:152620

Expression of disease reaction types in

soybean callus from resistant and susceptible

plants

AUTHOR(S):

Holliday, M. J.; Klarman, W. L.

CORPORATE SOURCE:

Dep. Bot., Univ. Maryland, College Park, MD, 20742,

SOURCE:

Phytopathology (1979), 69(6), 576-8

CODEN: PHYTAJ; ISSN: 0031-949X

DOCUMENT TYPE:

Journal

LANGUAGE:

English

Conditions influencing expression of disease reaction types were examd. in calluses derived from soybean plants resistant (cultivar Cutler 71) or susceptible (cultivar Cutler) to race 1 of Phytophthora megasperma sojae. Cutler 71 calluses were colonized less than those of Cutler when both were grown on medium contg. 6 or 10 mg of 2,4-D/L and incubated at 16 or 20.degree. prior to and following inoculation with P. megasperma sojae zoospores. Differences between colonization rates of Cutler and Cutler 71 calluses were greater in callus sections 5 mm thick than in thicker or thinner sections. Differences in colonization rates remained high with inoculum doses varying from 50 to 1000 zoospores per callus section. Sections of Cutler and Cutler 71 calluses 5 mm thick were colonized equally by race 3 of Phytophthora which is pathogenic to plants of both cultivars. No combinations of incubation temps., 2,4-D concns., sizes of calluses, or nos. of zoospores used for inoculum resulted in Cutler 71 calluses with the nearly abs. resistance to race 1 of ${\bf Phytophthora}$ found in whole plants of that cultivar.

94-75-7, biological studies IT RL: BIOL (Biological study)

(soybean calluses culture in media contg., colonization rates in relation to)

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

5-2 (Agrochemicals)

soybean cultivar callus Phytophthora; genotype soybean Phytophthora

IT Soybean

(disease reaction types in genotypes of)

IT Phytophthora megasperma sojae

(soybean cultivars susceptibility to, parameters of)

TT

(soybean resistance and susceptibility to Phytophthora megasperma sojae in relation to)

Plant tissue culture TT

(Phytophthora megasperma sojae susceptibility of soybean callus, genotypes in relation to)

94-75-7, biological studies IT

RL: BIOL (Biological study)

(soybean calluses culture in media contg., colonization rates in relation to)

L96 ANSWER 17 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1979:19203 HCAPLUS

DOCUMENT NUMBER:

90:19203

TITLE:

Modification of the disease resistance of

tobacco callus tissues by cytokinins

AUTHOR(S):

Haberlach, Geraldine T.; Budde, Allen D.; Sequeira,

Luis; Helgeson, John P.

CORPORATE SOURCE:

Dep. Plant Pathol., Univ. Wisconsin, Madison, WI, USA

SOURCE:

Plant Physiology (1978), 62(4), 522-5

CODEN: PLPHAY; ISSN: 0032-0889

DOCUMENT TYPE: LANGUAGE:

Journal English

AB The effects of differing cytokinin and auxin concns. on resistance of tobacco (Nicotiana tabacum) tissue cultures to race 0 of Phytophthora parasitica var nicotianae were examd. With 1 .mu.M 2,4-D tissues from resistant cultivars exhibited a "hypersensitive" reaction to zoospores of the fungus and subsequently were colonized only slightly. With susceptible cultivars or with tissues from resistant cultivars supplied with higher cytokinin levels (e.g.) mM kinetin), this hypersensitive reaction did not occur and tissues were heavily colonized. Benzylaminopurine and kinetin were particularly effective in eliminating both the hypersensitive reaction and disease resistance. Zeatin and 6-(3-methyl-2-butenylamino)purine were less effective. Increases in indoleacetic acid levels reversed the effects of high cytokinin concns. The balance of phytohormones apparently controls the host response to the fungus; thus, in this system, resistance or susceptibility can be studied without changing either host or fungal genotype.

IT 94-75-7, biological studies RL: BIOL (Biological study)

(tobacco callus disease resistance response to)

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

CC 11-5 (Plant Biochemistry)

ST tobacco disease resistance cytokinin; Phytophthora resistance tobacco cytokinin; auxin tobacco resistance Phytophthora

IT Tobacco

(disease resistance in callus tissues of, cytokinin
modification of)

IT Plant tissue culture

(disease resistance of, of tobacco, cytokinin modification of)

IT Phytophthora parasitica nicotianae

(tobacco callus tissue resistance to, modification by cytokinins)

IT Plant hormones and regulators
RL: BIOL (Biological study)

(auxins, modification of **disease** resistance in tobacco callus tissues by)

IT Plant hormones and regulators

RL: BIOL (Biological study)

(cytokinins, modification of disease resistance in tobacco

callus tissues by)

IT 87-51-4, biological studies **94-75-7**, biological studies 525-79-1 1214-39-7 1637-39-4 2365-40-4

RL: BIOL (Biological study)

(tobacco callus disease resistance response to)

L96 ANSWER 18 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1978:437966 HCAPLUS

DOCUMENT NUMBER:

89:37966

TITLE:

Activity of certain enzymes of phenol metabolism

during the herbicidal treatment of plants

AUTHOR(S):

Volynets, A. P.; Pal'chenko, L. A.

CORPORATE SOURCE:

Inst. Eksp. Bot., Minsk, USSR

PRYOR 09/781,695 Mater. Biokhim. Konf. Pribalt. Resp. B. SSR, 5th (1976 SOURCE:), Volume 2, 156-7. Editor(s): Sibul, I. K. Akad. Nauk Est. SSR: Tallinn, USSR. CODEN: 38BKAW Conference DOCUMENT TYPE: LANGUAGE: Russian GI OCH2CO2H I Soil application of 0.5 or 5 kg 2,4-D (I) [94-75-7], or 3.0 or 30 kg TCA [76-03-9]/ha stimulated phenylalanineammonia lyase [9024-28-6] and .beta.-glucosidase [9001-22-3], and raised the contents of flavone aglycons and flavonols in herbicideresistant lupine. In a herbicide-susceptible cultivar the increase in the phenylalanineammonia lyase activity and flavonoid content was less, and .beta.-glucosidase was inhibited by the herbicides. 94-75-7, biological studies RL: BIOL (Biological study) (lupine phenol metab. response to) RN 94-75-7 HCAPLUS Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) 0- CH2- CO2H 5-3 (Agrochemicals) lupine phenol metab dichlorophenoxyacetate TCA ST Lupine IT (2,4-D and TCA effect on phenol metab. of) IT Aglycons **Flavones** Flavonoids RL: BIOL (Biological study) (of lupine, 2,4-D and TCA effect on) IT Plant metabolism (of phenols, by lupine, 2,4-D and TCA effect on) IT **Flavones** RL: BIOL (Biological study) (hydroxy, of lupine, 2,4-D and TCA effect on) 76-03-9, biological studies 94-75-7, biological studies RL: BIOL (Biological study) (lupine phenol metab. response to) 9001-22-3 9024-28-6 IT RL: BIOL (Biological study) (of lupine, 2,4-D and TCA effect on) L96 ANSWER 19 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 1977:578664 HCAPLUS

DOCUMENT NUMBER:

87:178664

TITLE:

Metabolism of 2,4-dichlorophenoxyacetic acid by wheat

cell suspension cultures

AUTHOR(S):

Bristol, Douglas W.; Ghanuni, Ahmed Murad; Oleson,

Arland E.

CORPORATE SOURCE:

Dep. Biochem., North Dakota State Univ., Fargo, ND,

USA

SOURCE:

Journal of Agricultural and Food Chemistry (1977),

25(6), 1308-14 CODEN: JAFCAU; ISSN: 0021-8561

DOCUMENT TYPE: LANGUAGE:

Journal English

GI

Wheat cells in suspension culture absorbed 14C-labeled 2,4-D (I) [94-75-7] rapidly from B5 nutrient medium. After 4 days of incubation, the distribution of radiolabel in the culture system reached a steady state. After 6 to 8 days, the major metab. pathway for I involved ring hydroxylation followed by conjugation with sugars since over 37% of the applied radiolabel was present in the cells as H2O-sol./Et2O-insol. metabolites. Following hydrolysis, extraction into Et2O, and sepn. by TLC, 4-hydroxy-2,5-dichlorophenoxyacetic acid [2639-78-3] was identified as the major aglycone present. Lesser amts. of 4-hydroxy-2,3-dichlorophenoxyacetic acid [3004-84-0], 4-hydroxy-2chlorophenoxyacetic acid [7417-87-0], and I were detected. Et20-sol. amino acid conjugates and free I present in the cells after 6 to 8 days represented only 17 and 13%, resp., of the applied radioactivity. A considerable amt. (12%) of the applied radioactivity was bound to insol. ceklular tissue while 10% was present in the extracellular medium. Apparently 9.4% was lost from the system as a volatile metabolite. results of this model study are compared with those reported for other plants in tissue culture and support the hypothesis that the resistance of some plants to the herbicidal action of I is related to their species specific ability to accomplish detoxification by conversion to H2O-sol. metabolites.

IT 7417-87-0

RL: FORM (Formation, nonpreparative)

(formation of, from 2,4-D, by wheat in cell culture)

7417-87-0 HCAPLUS RN

Acetic acid, (2-chloro-4-hydroxyphenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

94-75-7, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(metab. of, by-wheat, in cell culture)

RN 94-75-7 HCAPLUS

CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

```
CC
     4-4 (Toxicology)
    wheat culture dichlorophenoxyacetate metab
ST
IT
    Wheat
        (2,4-D metab. by, in cell culture)
                3004-84-0 7417-87-0
IT
     2639-78-3
    RL: FORM (Formation, nonpreparative)
        (formation of, from 2,4-D, by wheat in cell culture)
    94-75-7, biological studies
IT
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (metab. of, by wheat, in cell culture)
L96 ANSWER 20 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1977:562591 HCAPLUS
DOCUMENT NUMBER:
                         87:162591
TITLE:
                         Effect of growth regulators on the development of
                         collar rot disease caused by the fungus
                         Phytophthora cactorum in apple trees
AUTHOR(S):
                         Plich M
CORPORATE SOURCE:
                         Res. Inst. Pomol., Skierniewice, Pol.
                         Fruit Science Reports (1976), 3(3), 33-42
SOURCE:
                         CODEN: FSREDB; ISSN: 0137-1479
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
    The auxins, IAA [87-51-4], NAA [86-87-3], and 2,4-D [94-75-7],
     BA [1214-39-7], and abscisic acid [21293-29-8] modified the susceptibility
   . of apple trees to P. cactorum, but the effects obtained were dependent on
     the lapse of time between the inoculation of the trees with the fungus and
     the growth regulator application. Auxin treatment 10 and 4 days before-
     infection attenuated disease development whereas BA greatly
     increased the size of necroses. These growth regulators had no influence
    when applied 4 days after infection. Gibberellicacid [77-06-5] did not
     produce a significant effect, but abscisic acid administered both before
     and after infection caused larger necroses. The effect of BA and abscisic
    acid were, however, dependent on the variety of apple as well as on the
     time of expt. performance. Growoth regulators active in vivo conditions
     showed relatively weak or no fungitoxic affects when applied in vitro.
     Growth regulators did not directly affect the development of the pathogen
     in the host tissues but they changed the susceptibility of apple trees by
    way of their influence on plant metab.
    94-75-7, biological studies
    RL: BIOL (Biological study)
        (apple collar rot disease response to)
RN
     94-75-7 HCAPLUS
    Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
C<sub>1</sub>
```

cc5-2 (Agrochemicals) growth regulator apple collar rot; Phytophthora apple ST plant hormone Plant hormones and regulators RL: BIOL (Biological study) (apple collar rot disease response to) IT Phytophthora cactorum (apple infestation with, plant hormones effect on) IT Apple (disease, collar rot, plant hormones effect on) 87-51-4, biological studies 94-75-7, IT 77-06-5 86-87-3 biological studies 1214-39-7 21293-29-8

RL: BIOL (Biological study) (apple collar rot disease response to)

L96 ANSWER 21 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1970:413407 HCAPLUS

DOCUMENT NUMBER:

73:13407

TITLE:

Effect of herbicides on isoflavone level in

lupine roots

AUTHOR(S):

Laman, M. A.; Valynets, A. P.; Mashtakoy, S. M.

Inst. Eksp. Bot., Minsk, USSR CORPORATE SOURCE:

SOURCE:

Vestsi Akademii Navuk BSSR, Seryya Biyalagichnykh

Navuk (1969), (6), 30-4 CODEN: VABBA3; ISSN: 0002-3558

DOCUMENT TYPE:

Journal Belorussian

LANGUAGE:

Yellow lupine plants of the varieties Belorusskii (A) and

Baravlyanskii (B) were treated with solns. of Na trichloroacetate, dalapon, and the Na salt of 2,4-D that were poured on the soil surface. One, 5, and 11 days after application of the herbicides, samples of the

plant roots were treated with steam, dried at 50-60.degree., and on grinding extd. with ether and then with 80% EtOH. The extd. material was sepd. by two-dimensional paper chromatog. by using iso-BuOH-AcOH-H2O (4:1:5) and 5% AcOH as solvents. Spots corresponding to 5 substances (1-5) were obtained. These substances, were identified as isoflavone polyphenols. The high soly. of 1-5 in solvent mixts. contg. H2O indicated that the isoflavone derivs. were present in the form of glycosides. The uv spectra showed that an OH group in position 7 was substituted with the sugar residue, while an OH group in position 5 was free. The content of isoflavones 1-5 in the roots depended on the length of time during which the herbicide acted and the resistance of the lupine variety to herbicides. Whereas in the resistant variety A the content of 1-5 24 hr after application of Cl3CCO2Na and dalapon decreased by 7 and 11.4%, resp., vs. untreated controls, it increased by 16 and 32%, resp., in variety B. Apparently, 1-5 act in the roots of lupines as antagonists of indolylacetic acid.

IT 3653-48-3

> RL: BIOL (Biological study) (isoflavone formation by roots in response to, soil

treatment with)

RN 3653-48-3 HCAPLUS

CN Acetic acid, (4-chloro-2-methylphenoxy)-, sodium salt (9CI) (CA INDEX NAME)

18 (Plant Growth Regulators)

isoflavones lupine herbicides; lupine isoflavones herbicides; herbicides lupine isoflavones; roots lupine isoflavones; dalapon lupine isoflavones; chloroacetate lupine is flavones

IT Soils

(isoflav ne formation in r ts in herbicide-treated)

IT Lupines

```
(isoflavones from r ots of, herbicide effect on)
     Isoflavone, derivs.
     RL: FORM (Formation, nonpreparative)
        (formation of, by roots in herbicide-treated soils)
IT
     650-51-1
     RL: BIOL (Biological study)
        (isoflavone formation by roots in response to soil
        treatment with)
     75-99-0 3653-48-3
ΙT
     RL: BIOL (Biological study)
        (isoflavone formation by roots in response to, soil
        treatment with)
L96 ANSWER 22 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                         1956:61867 HCAPLUS
DOCUMENT NUMBER:
                         50:61867
ORIGINAL REFERENCE NO.:
                         50:11593h-i
TITLE:
                         Downy mildew of onions; results of further spray trial
AUTHOR(S):
                         <del>Doepel, R. F.</del>
SOURCE:
                         J. Dept. Agr. W. Australia (1956), 5, 185-6,189-90
DOCUMENT TYPE:
LANGUAGE:
                         Unavailable
     cf. C.A. 50, 2909b.
                          The effectiveness of zineb fungicide sprays for
     controlling downy mildew of onions was confirmed by further trials in
    which zineb (1 1/2 lb./100 gal.) was compared with Cu
     oxychloride (3 1/3 lb./100 gal.). Plots sprayed with zineb yielded 20%
     more marketable onions than the Cu oxychloride and control
     plots, and the disease was also greatly reduced in seed
     crops. Use of Triton B1956 spreader (6 fluid oz./100 gal.)
     improved the coverage of plants.
     94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
IT
        (in mushroom-mildew control)
     94-75-7 HCAPLUS
RN
     Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
           0-CH2-CO2H
     15A (Pesticides and Crop-Control Agents)
IT
     Peronosporaceae
        (control of, on onions)
IT
     Onions
        (downy-mildew control on)
     Fungicides or Fungistats
IT
        (for mildew, on onions)
IT
    Roccal
        (in mushroom-mildew control)
TT
    Copper chlorides
        (basic, in downy-mildew control on onions)
     Carbamic acid, ethylenebis[dithio-, disodium and Zn salts
TT
     Carbamic acid, methyldithio-, sodium salt
     Glutarimide, 3-[2-(3,5-dimethyl-2-oxocyclohexyl-2-hydroxyethyl]-
        (Actidione)
    Hydrazine, sulfate, Cu complex
        (in mushroom-mildew control)
     Carbamic acid, ethylenebis[dithio-, zinc salt
IT
        (in onion-downy-mildew control)
IT
     14798-03-9, Ammonium
        (compds., substituted, alkylbenzyldimethyl-chlorides, in
        mushroom-mildew control)
ΙT
     148-24-3, 8-Quinolinol
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(copper complexes, in mushroom-mildew control) 50-00-0, Formaldehyde 51-28-5, Phenol, 2,4-dinitro-57-13-6, Urea 79-57-2, Oxytetracycline 82-68-8, Benzene, pentachloronitro-87-86-5. Phenol, pentachloro- 93-23-2, Isoquinolinium, 2-dodecyl-, bromide 94-75-7, Acetic acid, (2,4-dichlorophenoxy)- 100-97-0, Hexamethylenetetramine 101-05-3, s-Triazine, 2,4-dichloro-6-o-117-18-0, Benzene, 1,2,4,5-tetrachloro-3-nitrochloroanilino-118-74-1, Benzene, hexachloro-123-33-1, 3,6-Pyridazinedione, 1,2-dihydro- 124-40-3, Dimethylamine 128-04-1, Carbamic acid, dimethyldithio-, sodium salt 131-52-2, Phenol, pentachloro-, sodium dimethyldithio-, sodium salt 132-27-4, Phenol, o-phenyl-, sodium deriv. 1319-77-3, Cresol 1336-21-6, Ammonium hydroxide 1403-61-8, Fradicin 2275-75-4, N 1045 2492-26-4, 2312-76-7, o-Cresol, 4,6-dinitro-, sodium deriv. Benzothiazole, 2-mercapto-, sodium deriv. 7778-54-3, Calcium hypochlorite 17273-33-5, 2-Naphthaleneacetic acid, sodium salt (in mushroom-mildew control)

L96 ANSWER 23 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

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ACCESSION NUMBER:
                           1955:75150 HCAPLUS
DOCUMENT NUMBER:
                           49:75150
ORIGINAL REFERENCE NO.:
                          49:14253i,14254c-g
                           Chemotherapeutic application of some compounds to rice
TITLE:
                           plants and the outbreak of Helminthosporium
                           Teaf spot
                           Akai, Shigeyasu
AUTHOR(S):
CORPORATE SOURCE:
                           Kyoto Univ.
                           Shokubutsu Byogai Kenkyu (1955), 5, 45-56
SOURCE:
                           CODEN: SBYKAH; ISSN: 0370-8845
DOCUMENT TYPE:
                           Journal
                           Enalish
LANGUAGE:
     When rice seed was soaked in 2 p.p.m. CuSO4 soln. for 24 hrs. at room
     temp., no remarkable effect was shown in the controlling of
     diseased spots on mature leaves, but when rice seedlings
     were treated with the soln. of 2 p.p.m. CuSO4 and 200 p.p.m. boric acid,
     an appreciable effect was shown. When rice seed and roots of young seedlings were treated with 1-naphthaleneacetic acid and
    3-indoleacetic acid-solns. (10-100 p.p.m.), a difference in their effects was noticed for the root treatment but not for the seed one.

When roce seed was soaked in-solns. of (2,4-D), 10-3, 10-4, 10-5, 10-6%, and
     a control soln., for 48 hrs. at 36.degree., the percentages of
     diseased spots were, resp., 31, 41, 66, 62, and 100%. In this
     expt., the growth of the seedlings was delayed markedly when soaked in a
     10-3% soln., but afterwards they recovered and almost no difference in the
     growth was obtained between plots tested. When rice seedlings were
     treated twice with 3 naphthoic acid derivs., 1,4-dihydronaphthoic (I)
     1,2,3,4-tetrahydronaphthoic (II), and "2,4,5-trihydronaphthoic acid"
     (III), in the concns. of 2 p.p.m. at first and 5 p.p.m. after 4 days, the
     percentages of diseased spots were, resp., 21, 11, and 61%. I
     inhibited 50% germination of conidia in 10-2% concn., II did not inhibit
     in any concn., and III had a considerable fungicidal effect, permitting
     only 7% germination of conidia in 2 .times. 10-2% concn. I and II had no
     phytotoxic effect on rice plants, but III gave some injury. The
     chemotherapeutic action of vitamin K3 (2-methyl-1,4-naphthoquinone) was
     studied by soaking seed and roots of seedlings in 0.01-0.02%
     solns. At first the plants showed a vigorous growth, but the
     difference in plant growth between the treated plants
     and control diminished with the lapse of time. A significant protection
     from the disease was obtained in the treated plants by
     soaking in vitamin K3 soln., and the enlargement of diseased
     spots on leaves of treated plant was more delayed than
     that of the control. Pentachlorophenol, its Na salt, and
     pentachlorophenoxyacetic acid reduced markedly the infection by this
   fungus. However they caused injury in 0.0025% soln. to rice
     plants, changing the color of the leaves to yellow.
     94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
         (in Helminthosporium control on rice)
     94-75-7 HCAPLUS
RN
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CN Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)

CC 15A (Pesticides and Crop-Control Agents)

IT Helminthosporium

(control on rice, and plant regulators in relation thereto)

IT Fungicides or Fungistats

(for Helminthosporium on rice)

IT Plant regulators

(in Helminthosporium control on rice)

IT <u>Rice</u>

(Helminthosporium control on, and plant regulators in relation thereto)

IT 3-Indoleacetic acid

Boric acid

(in Helminthosporium control on rice)

IT 13295-81-3, Propionic acid, 3-chloro-, 5-nitrofurfuryl ester

(in Coccidioides immitis control)

L96 ANSWER 24 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1950:57997 HCAPLUS

DOCUMENT NUMBER:

44:57997

ORIGINAL REFERENCE NO.:

44:10992a-e

TITLE:

Gladiolus research in Florida, 1949-1950 season

AUTHOR(S):

Magie, Robert O.

CORPORATE SOURCE:

Florida Agr. Expt. Sta., Bradenton

SOURCE:

North American Gladiolus Council Bulletin (1950), No.

23, 81-2,84,87

CODEN: NOGCA7; ISSN: 0029-2370

DOCUMENT TYPE:

Journal

LANGUAGE:

Unavailable

cf. C.A. 44, 5511b. The weather during the past season was discussed in relation to the development of Curvularia and Stemphylium diseases and Botrytis sclerotia. Several new virus diseases and harvesting and storage problems are being investigated. In Florida, a great deal of trouble is encountered owing to overfertilization. The importance of including B, Fe, and Mn salts in fertilizer is pointed out; deficiencies may not show up until the following year. The "running out" of older corm stocks may be due to this deficiency. For pre-emergence control of weeds, the insol. 2,4-D was used in a wettable form, at 3 oz. per 1000 ft. of row in an 18-in. band centering on the row. (This form was used to prevent root injury due to excessive rainfall.) While results were satisfactory, the material was difficult to maintain in suspension. The most effective treatment was a spray mixt. of 2,4-D and Aero-Cyanate applied when the weeds were about 1/2 in. high and when the gladiolus leaves from No. 3 corms had 2-3 leaves. Rate of application was 5 lb. of Aero-Cyanate and 2 1/2 lb. of 2,4-D in 60 gal. per acre of rows spaced 3 ft. apart, half the area being sprayed. Aero-Cyanate alone was effective over a shorter period, when applied at 5-10 lb. per acre. Spraying must be carried out when the weeds are small, and repeated as necessary. Phenyl Hg acetate was effective and apparently quite safe for weed control at 3 lb. per acre, but was too expensive. Shell Oil 130 (pentachlorophenol) was discontinued

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because of the danger to the operator's eyes. Methyl bromide is
      recommended for fumigating seed beds and beds for cormel planting to
      control nematodes, weeds, and soil fungi.
      94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
(in weed control, in gladiolus)
 IT
 RN
      94-75-7 HCAPLUS
      Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
 CN
            0- CH2- CO2H
CC
      15A (Economic Poisons)
      Nutrition, plant
         (boron, Fe and Mn in, of gladiolus)
 IT
      Fusarium
         (control of, on gladiolus)
 TT
      Curvularia
      Sclerotinia sclerotiorum
         (control on gladiolus)
 IT
      Gladiolus
         (diseases of)
 IT
      Fertilizers
         (gladiolus diseases and)
 TT
      Weed control
         (in gladiolus)
 IT
      Fumigation
         (of gladiolus with MeBr)
 IT
      Stemphylium
         (on gladiolus and its control)
 IT
      Gladiolus
         (Fusarium control on corms)
 IT
      74-83-9, Methane, bromo-
         (as gladiolus fumigant)
                       7439-96-5, Manganese
                                               7440-42-8, Boron
 IT
      7439-89-6, Iron
         (in gladiolus nutrition)
 IT
      62-38-4, Mercury, phenyl-, acetate
         (in weed control in gladiolus)
· IT
      87-86-5, Phenol, pentachloro- 94-75-7, Acetic acid,
      (2,4-dichlorophenoxy)- 590-28-3, Potassium cyanate
         (in weed control, in gladiolus)
 L96 ANSWER 25 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
 ACCESSION NUMBER:
                           1949:52190 HCAPLUS
 DOCUMENT NUMBER:
                           43:52190
 ORIGINAL REFERENCE NO.:
                          43:9338e-g
                           Controlling carnation diseases
 TITLE:
 AUTHOR(S):
                          Matheron, E. John
                          N. Y. State Flower Growers Bull. (1949), No. 49, 11-12
 SOURCE:
 DOCUMENT - TYPE:
                          Journal
                          Unavailable-
 LANGUAGE:
      As a footing agent, 1 g. of indolebutyric acid is mixed with 2 lb. of a
      very refined face powder; this powder does not permit too much to adhere
      to the bottom of the cutting. Cuttings are rooted in sand, which has been
      generously dusted with Fermate; the sand is changed for each batch of
      cuttings. Most of the rooted cuttings are benched where they will be
      undisturbed until final planting or benching time; too frequent
      transplanting tends to spread disease. All young plants
      that are placed indoors are dipped in a suspension of 2 lb. Fermate/100
      gal. of water. Spiders are controlled with 15% wettable Parathion powder
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which is sometimes combined with the Fermate. Tech. Parathion has also been used at the rate of 5 oz./100 gal. of water; in each case 6 oz. of a

teaspoons/gal. of water has given promising results. Limited trials with Fulex soil treatments gave neg. results. 94-75-7, Acetic acid, (2,4-dichlorophenoxy)-IT (in weed control) RN 94-75-7 HCAPLUS Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME) CN - CH2- CO2H 15A (Economic Poisons) Grasses (control of Bermuda and Johnson) IT Spiders (control of, on carnations) IT Carnation (Dianthus) (cutting, propagation and disease control) IT (disinfection, for carnations with Fulex) IT Insecticides (for spiders on carnations) IT Plant regulators (indolebutyric acid as, for carnation cuttings) IT 3-Indolebutyric acid (as carnation-rooting agent) Carbamic acid, dimethyldithio-, iron salt IT (in carnation-disease control) IT 56-38-2, Parathion (in carnation spider control) IT 117-80-6, Dichlone 3689-24-5, Fulex (in carnation-disease control) IT 94-75-7, Acetic acid, (2,4-dichlorophenoxy)-(in weed control) L96 ANSWER 26 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN ACCESSION NUMBER: 1949:52066 HCAPLUS DOCUMENT NUMBER: 43:52066 ORIGINAL REFERENCE NO.: 43:9313a-d Science in the land [Annual Rept.] TITLE: AUTHOR(S): Martin, W. H. SOURCE: N. Jersey Agr. Expt. Sta., Ann. Rept. (1947), Volume Date 1945-1946 7-109 DOCUMENT TYPE: Journal Unavailable LANGUAGE: cf. C.A. 41, 1774e. Progress reports are made, of which the following are of chem. interest: the effects of feeding thyroprotein to dairy cattle, effects on Ca and P metabolism when traces of Mn, Cu, and Fe are added to the ratio of milking cows, use of thiouracil in swine fattening, phosphoric acid deficiencies in feeding, fertilization of bromegrass; effects of 2,4-D on hedge and field bindweed, mustard, watercress, horsenettle, curled dock, and dandelions; control of late blight of potatoes, relationship between boron and Ca in the nutrition of the tomato plant, fungicides for tomatoes, methylcellulose and Tersan (tetramethylthiuram disulfide) for the control of onion smut, control of European corn borer on sweet corn, effect of weather on the sugar and acid content of peaches, p-dichlorobenzene for the control of peach tree borer, fungicides for apple diseases, injurious

effects of industrial fumes upon green plants in New Jersey,

on greenhouse plants, relationship between light intensity and

fertilization of strawberries, azobenzene for the control of red spiders

spreader-sticker is added per 100 gal. of water. Phygon at 2

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absorption of nutrients by plants, control of stink worm with benzene hexachloride, control of poison ivy with borax and ammonium sulfamate, fertilization requirements of tomatoes and corn, Mg deficiencies of blueberries, corn, potatoes, sweet potatoes, and tomatoes, relationship between ir n and Mn needs of plants, effect of factory fumes upon the soil, feeding of Vitamin D to oysters, rotenone content of devil's shoestring (Tephrosia virginiana) grown in New Jersey, fire-resistant barns made with cement and asbestos, protection of silo walls against the action of juices, and nutritive value of the protein of green snap beans.

94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
(in weed control)

94-75-7 HCAPLUS
Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
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IT

RN

CN

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C1
           0-CH2-CO2H
     Ċ1
CC
     15 (Soils and Fertilizers)
IT
     Agriculture (includes Agronomy)
        (New Jersey Agr. Expt. Station report on)
IT
     Fungicides or Fungistats
        (New Jersey Expt. Sta. reports on)
IT
     Feeding experiments
        (New Jersey Expt. Station reports on)
IT
     Peaches
        (acids and sugars in, weather effect on)
TT
     Red spider mite
        (azobenzene in control of)
IT
     Fire-resistant materials
        (barns)
IT
     Nutrition, animal
        (bean (green-snap) protein in)
IT
        (blight (late) control on)
IT
     Grasses
        (brome, fertilizer expts. with)
     Absorption, biological
IT
        (by plants, light intensity and)
IT
     Poison ivy
     Pyrausta nubilalis and(or) European corn borer
        (control of)
IT
     Dock
        (control of curled, 2,4-D in)
     Bindweed
     Dandelion and(or) Taraxacum officinale
     Mustard
     Solanum carolinense
     Water cress
        (control of, 2,4-D in)
ΙT
     Sanninoidea exitiosa and(or) Peachtree borer
        (control of, p-dichlorobenzene in)
IT
    Earthworms
        (control with hexachlorocyclohexane)
IT
     Thyroproteins
        (effect on dairy cattle)
IT
     Fumes
     Fumes
        (effect on green plants and soil)
IT
     Insecticides
        (expt. sta. reports on)
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Fertilizers
IT
        (expts., in New Jersey)
IT
     Soils
        (factory-fume effect on)
ΙT
     Corn
     Strawberries
     Tomatoes
        (fertilizer expts. with)
IT
     Buildings
        (fire-resistant barn-type)
IT
     Plants
        (fume (industrial-) effect on)
IT
     Apples
        (fungicides for)
    Nutrition, plant
(in New Jersey)
IT
IT
     Acids
        (in peaches, effect of weather on)
IT
     Sugars
        (in peaches, weather effect on)
IT
     Light
        (intensity of, absorption of nutrients by plants and)
IT
     Blueberries
     Corn
     Potatoes
     Sweet potatoes
     Tomatoes
        (magnesium deficiency in)
IT
     Tomatoes
        (nutrition and pest control on)
IT
     Proteins
        (of beans, nutritive value of)
IT
    Metabolism, animal
        (of calcium and P, by milking cows, effect of Cu, Fe
        and Mn on)
IT
    Nutrition, plant
        (of tomatoes, relation between B and Ca in)
IT
     Silos
        (protection of walls of)
     Beans and(or) Phaseolus
IT
        (proteins of green snap, nutritive value of)
     Tephrosia virginiana
IT
        (rotenone in)
IT
     Onions
        (smut control on)
IT
    Milk
        (thyroprotein in feeding and)
IT
     Oysters
        (vitamin D effect on)
IT
     Weed control
        (with 2,4-D)
     Cellulose, methyl ether
IT
        (in onion-smut control)
     Vitamin, D (antirachitic)
IT
        (oyster feeding with)
IT
     7664-38-2, Phosphoric acid
        (deficiencies in feeding)
     7439-95-4, Magnesium
IT
        (deficiency of, in plants in New Jersey)
                       7439-96-5, Manganese
                                               7440-50-8,
IT
     7439-89-6, Iron
     Copper
        (effect on metabolism of Ca and P during lactation in cattle)
IT
     83-79-4. Rotenone
        (in devil's shoestring)
     137-26-8, Disulfide, bis(dimethylthiocarbamoyl)
IT
        (in onion-smut control)
IT
     106-46-7, Benzene, p-dichloro-
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(in peach-tree borer control)
IT
     7439-96-5, Manganese
        (in plant nutrition, Fe and)
IT
     7439-89-6, Iron
        (in plant nutrition, Mn and)
     1303-96-4, Borax 7773-06-0, Ammonium sulfamate
IT
        (in poison-ivy control)
     103-33-3, Azobenzene
TT
        (in red-spider control on greenhouse plants)
     608-73-1, Cyclohexane, 1,2,3,4,5,6-hexachloro-
IT
        (in stink-worm control)
IT
     141-90-2, Uracil, 2-thio-
        (in swine fattening)
IT
     7440-70-2, Calcium
        (in tomato nutrition, B and)
IT
     7440-42-8, Boron
        (in tomato nutrition, Ca and)
     94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
        (in-weed control)
                          7723-14-0. Phosphorus
IT
    7440-70-2, Calcium
        (metabolism of, by cows, effect of Cu, Fe and Mn
L96 ANSWER 27 OF 27 HCAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                          1947:8535 HCAPLUS
DOCUMENT NUMBER:
                          41:8535
ORIGINAL REFERENCE NO.:
                          41:1791f-i,1792c-d
TITLE:
                          Research in agriculture (Annual report, 1944-45)
                          Taggart, W. G.
AUTHOR(S):
CORPORATE SOURCE:
                          Baton Rouge
SOURCE:
                          Louisiana Agr. Expt. Sta., Ann. Rept. (1945), Volume
                          Date 1944-1945 143 pp.
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          Unavailable
     Research studies on the following subjects are briefly summarized:
     ascorbic acid value of tomatoes canned by 5 home-processing methods;
     varietal differences in the ascorbic acid value of strawberries; carotene
     and ascorbic acid content of sweet potatoes; vitamin A content of milk and
     butter; enrichment of rice; growth stimulants for microbiol. biotin assay;
     nutritional status of pregnant women; utilization of ascorbic acid of
     leafy vegetables by humans; exptl. lathyrism; detoxication of tung meal;
     toxic principles of the tung nut; freezing strawberries, shrimp, French
     fried potatoes, and freezing peaches with added ascorbic acid; dehydration
     of sweet potatoes; effect of org. matter, plowing, and vegetable cover on
     runoff; soybean meal and peanut meal as protein supplements for fattening
     pigs; effects of depth of application on the loss of N from flooded soil;
     sugar-cane fertilization; fertilization and rotation for cotton; oat
     fertilization; mineral deficiencies of La. dairy herds; producing pasture
     with com. fertilizer and manure; DDT-nicotine combination effective
     against cabbage worms; chem. control of sand wireworm; controlling
     velvet-bean caterpillar on soybeans with DDT or cryolite; nicotine in dry
     concentrate form for cotton aphid; sugar-cane borer control; Na
     fluosilicate more toxic than cryolite to the borer; nicotine and
     nicotine-rotenone dust mixts. for control of turnip aphid; onion thrip
     control; control of cockroaches, fleas, brown dog tick, and flies; seasonal changes in carotene content of sweet potatoes; waxing stored
     sweet potatoes; control of nematodes by soil treatment; fermate controls
     mildew and anthracnose diseases of cucumbers; 2,4-D on various
    weeds; soil rot of sweet potatoes controlled by applying S; soil treatment for control of shallot white rot; physiology of the avian thyroid;
     coccidiosis control; Johne's disease in cattle;
     gastro-intestinal nematode parasites of cattle; Crotalaria spectabilis
     poisoning in Louisiana livestock; drugs for the control of pinkeye in
     cattle; effect of lime on production of strawberries; green feeds for
     hens; P fertilization with various carriers; effect of straw with and
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without fertilizers for rice; residual effect of Ca arsenate on rice yields; substitutes, adjuvants, and reduced dosages for rotenone and

PRYOR 09/781,695

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pyrethrum for control of insects attacking cole cr ps; and chem.
     and microbiol. studies of soil from wilt-free and wilt-infested areas.
     94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
IT
     (in weed control)
94-75-7 HCAPLUS
RN
     Acetic acid, (2,4-dichlorophenoxy)- (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
           0-CH2-CO2H
     15A (Economic Poisons)
     Lathyrism
IT
IT
     Sweet potatoes
        (Expt. Sta. reports on, of Louisiana)
IT
     Frozen foods
        (French-fried potatoes, peaches, shrimp and strawberries)
IT
     Carotene
     Nutrition, animal
     Strawberries
        (Louisiana Expt. Sta. report on)
IT
     Insecticides
     Soils
        (Louisiana Expt. Sta. reports on)
IT
     Pyrethrum
        (as cole-crop insecticide)
     Soybean meal
IT
        (as protein supplement, for fattening hogs)
IT
     Poisoning and Intoxication
        (by Crotalaria spectabilis, of livestock)
     Cabbage
TT
        (caterpillar control on, DDT-nicotine combinations in)
     Conjunctivitis
IT
        (control in cattle)
     Aphis gossypii
IT
     Coccidiosis
     Cockroaches
     Diatraea saccharalis and(or) Sugarcane borer
     Fleas
     Flies
     Horistonotus uhlerii and(or) Sand wireworm
     Rhipicephalus sanguineus and(or) Brown dog tick
     Rhopalosiphum pseudobrassicae and(or) Turnip aphid
     Thrips tabaci and(or) Onion thrips
        (control of)
IT
     Cucumber
        (control of anthracnose and mildew on)
IT
     Anticarsia gemmatilis and (or) Velvetbean caterpillar
        (control of on soybeans)
IT
     Nematodes
        (control of, in cattle and soils)
IT
     Tung meal
        (detoxication of)
IT
     Straw
        (effect on rice)
IT
     Rice
     Rice
        (enrichment of, and effect of Ca arsenate residues, fertilizers and
        straw on)
IT
     Fertilizers
        (expts., in Louisiana)
IT
     Cotton
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(fertilization and rotation for)
IT
     0ats
     Pasture
     Sugar cane
        (fertilizer expts. with)
IT
     Growth substances
        (for biotin microbiol. assay)
IT
     Peaches
     Shrimp
        (freezing of)
IT
     Potatoes
        (french-fried, freezing of)
     Feeding stuffs
IT
        (green, for poultry)
IT
     Dysentery
        (in cattle)
     Peanut meal
IT
        (in fattening hogs)
IT
     Poisons
        (in tung nut)
IT
     Conjunctivitis
        (infectious, pharmaceuticals for)
IT
     Brassica
        (insect control on)
IT
     Crotalaria spectabilis
        (livestock poisoning by)
     Pregnancy
IT
        (nutrition in)
IT
     Waxing
        (of potatoes (sweet))
IT
     Drying
        (of sweet potatoes)
IT
     Detoxication
        (of tung meal)
IT
     Thyroid gland
        (physiology of avian)
ΙT
     Vitamins
        (rice enrichment with)
IT
     Lime
        (strawberry production and)
     Aleurites
IT
        (toxic principles of)
IT
     Soybeans
        (velvet-bean-caterpillar control on)
IT
     Butter
     Milk
        (vitamin A in)
IT
     Tomatoes
        (vitamin C in canned)
IT
     Canned goods
        (vitamin C in tomato)
IT
     Vegetables
        (vitamin C utilization from leafy, by humans)
IT
     Shallots
        (white rot of, control of)
IT
     Weed control
        (with 2,4-D)
     Feeding experiments
IT
        (with peanut meal and soybean meal on hogs)
IT
     Vitamin, A
        (in butter and milk)
     Carbamic acid, dimethyldithio-, iron salt
IT
        (in control of anthracnose and mildew on cucumbers)
     50-81-7, Vitamin, C
IT
        (Louisiana Expt. Sta. report on)
     15096-52-3, Cryolite
ΙT
        (as insecticide, in La.)
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IT
      54-11-5, Nicotine
         (as insecticide, in Louisiana)
ΙT
      58-85-5, Biotin
     (detn. of, growth stimulants for microbiol.) 7778-44-1, Calcium arsenate
IT
         (effect on rice)
     16893-85-9, Sodium fluosilicate
IŢ
         (in sugar-cane-borer control)
IT
     7704-34-9, Sulfur
     (in sweet-potato-soil-rot control)
94-75-7, Acetic acid, (2,4-dichlorophenoxy)-
ΙT
         (in weed control)
     83-79-4, Rotenone
IT
         (insecticide from)
      50-29-3, Ethane, 1,1,1-trichloro-2,2-bis(p-chlorophenyl)-
IT
         (reviews on)
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